

GenCore version 5.1.3  
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OM nucleic - nucleic search, using sw model

Run on: January 21, 2003, 14:54:52 : Search time 159.389 Seconds  
(without alignments)  
268.450 Million cell updates/sec

Title: US-09-853-688-36

Perfect score: 19

Sequence: 1 ttaggaagctcgggtgc 19

Scoring table: IDENTITY NUC

Gapop 10.0 , Gapext 1.0

Searched: 2185239 seqs, 1125999159 residues

Total number of hits satisfying chosen parameters: 4370478

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

N\_Geneseq\_101002:\*

- 1: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1980.DAT:\*
- 2: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1981.DAT:\*
- 3: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1982.DAT:\*
- 4: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1983.DAT:\*
- 5: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1984.DAT:\*
- 6: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1985.DAT:\*
- 7: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1986.DAT:\*
- 8: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1987.DAT:\*
- 9: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1988.DAT:\*
- 10: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1989.DAT:\*
- 11: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1990.DAT:\*
- 12: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1991.DAT:\*
- 13: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1992.DAT:\*
- 14: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1993.DAT:\*
- 15: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1994.DAT:\*
- 16: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1995.DAT:\*
- 17: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1996.DAT:\*
- 18: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1997.DAT:\*
- 19: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1998.DAT:\*
- 20: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1999.DAT:\*
- 21: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA2000.DAT:\*
- 22: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA2001A.DAT:\*
- 23: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA2001B.DAT:\*
- 24: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA2002.DAT:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

# SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	19	100.0	19	AAS18883	Growth hormone 1 g
c	19	100.0	3700	AAS18886	Growth hormone 1 g
3	16.4	86.3	5333	AAS46366	Tumour suppressor
4	16.4	86.3	5333	ABL32783	Human immune syste
c	15.8	83.2	65	ABN31023	Rat spliced transc
c	15.8	83.2	428	AAK54271	Murine transport a
7	15.8	83.2	730	AAF08345	Fusarium venenat
c	15.8	83.2	1612	AAAD11871	Wild-type sheep BM
c	15.8	83.2	1612	AAAD11872	Booroola sheep mut

c	10	15.8	83.2	2246	16	AAQ74064	Rat serotonin-1c c
c	11	15.8	83.2	2246	19	AAV48290	Rat 5-HT2C seroton
c	12	15.8	83.2	2246	19	AAV48291	Rat 5-HT2C S312F m
c	13	15.8	83.2	2246	19	AAV48298	Rat 5-HT2C seroton
c	14	15.8	83.2	2246	19	AAV48299	Rat 5-HT2C S312K m
c	15	15.8	83.2	2246	19	AAH48279	Rat 5-HT2C seroton
c	16	15.8	83.2	6109	23	AAH78667	Murine Col5a3 cDNA
c	17	15.8	83.2	10329	21	AAZ50264	Genomic DNA of pot
c	18	15.8	83.2	10329	21	AAZ50653	Genomic DNA encodi
c	19	15.4	81.1	170	24	ABL77410	Human ovarian canc
c	20	15.4	81.1	456	22	ABA58575	Human foetal liver
c	21	15.4	81.1	456	22	AAK06696	Human brain expres
c	22	15.4	81.1	456	22	AAK32395	Human bone marrow
c	23	15.4	81.1	456	22	AAI38237	Probe #6923 used t
c	24	15.4	81.1	456	24	ABS07178	Human genome-deriv
c	25	15.4	81.1	700	22	AAH92715	Human inflammatory
c	26	15.4	81.1	881	24	ABO46242	Oligonucleotide fo
c	27	15.4	81.1	881	24	ABO46243	Oligonucleotide fo
c	28	15.4	81.1	901	24	ABO33436	Oligonucleotide fo
c	29	15.4	81.1	901	24	ABO33437	Oligonucleotide fo
c	30	15.4	81.1	1343	21	AAZ59837	Human secreted pro
c	31	15.4	81.1	4038	10	AAH90382	Genes encoding hum
c	32	15.4	81.1	34917	22	AAK70686	Human immune/haema
c	33	15	78.9	392	24	ABK62503	Rat sequence diffe
c	34	15	78.9	2207	24	ABK63542	Rat sequence diffe
c	35	15	78.9	24757	22	AAZ27687	DNA encoding novel
c	36	15	78.9	24757	22	AAZ33481	DNA encoding human
c	37	14.8	77.9	393	21	AAFL5743	Human prostate can
c	38	14.8	77.9	393	22	AAK00856	Human CDNA clone H
c	39	14.8	77.9	558	22	AAK36046	Human bone marrow
c	40	14.8	77.9	558	22	AAI41762	Probe #10448 used
c	41	14.8	77.9	640	22	AAH71209	Human cervical can
c	42	14.8	77.9	653	21	AAZ75146	Human ORFX ORF701
c	43	14.8	77.9	653	24	ABN25315	Human ORFX polynuc
c	44	14.8	77.9	698	22	AAZ25821	C. glutamicum quan
c	45	14.8	77.9	787	23	AAZ71701	DNA encoding novel

## ALIGNMENTS

RESULT 1  
AAS18883  
ID AAS18883 standard; DNA; 19 BP.  
XX AAS18883;  
AC AAS18883;  
XX  
XX 12-MAR-2002 (first entry)  
XX Growth hormone 1 gene (GH1) specific fragment, PCR primer GH1R.  
XX  
XX Growth hormone 1; GH1; osteopathic; gene therapy; protein therapy;  
KW diabetes; obesity; infection; acromegaly; gigantism; sodium retention;  
KW water retention; metabolic syndrome; mood disorder; sleep disorder;  
KW Growth hormone dysfunction; familial growth hormone deficiency;  
KW short stature; pituitary storage defect; human; PCR primer: GH1R; ss.  
XX Homo sapiens.  
XX  
XX WO200185993-A2.  
XX  
XX 15-NOV-2001.  
XX  
XX 14-MAY-2001; 2001WO-GB02126.  
XX  
XX 12-MAY-2000; 2000GB-0011459.  
XX 14-JUL-2000; 2000EP-0306004.  
XX  
XX (UTWA-) UNIV WALES COLLEGE OF MEDICINE.  
XX  
XX Cooper DN, Procter AM, Gregory J, Millar DS;  
XX WPT; 2002-089798/12.  
XX

XX	FT	variation	replace(128,T)	
PT	FT	variation	/*tag= d	"Single nucleotide polymorphism"
PT	FT	variation	/standard_name=	replace(134,G)
PT	FT	variation	/*tag= e	"Single nucleotide polymorphism"
XX	FT	variation	/standard_name=	replace(135,T)
XX	FT	variation	/*tag= f	"Single nucleotide polymorphism"
XX	FT	variation	/standard_name=	replace(135,C)
CC	FT	variation	/*tag= g	"Single nucleotide polymorphism"
CC	FT	variation	/standard_name=	replace(136,G)
CC	FT	variation	/*tag= h	"Single nucleotide polymorphism"
CC	FT	variation	/standard_name=	replace(141,G)
CC	FT	variation	/*tag= i	"Single nucleotide polymorphism"
CC	FT	variation	/standard_name=	replace(179,C)
CC	FT	variation	/*tag= j	"Single nucleotide polymorphism"
CC	FT	variation	/standard_name=	replace(188,T)
CC	FT	variation	/*tag= k	"Single nucleotide polymorphism"
CC	FT	variation	/standard_name=	replace(218,A)
CC	FT	variation	/*tag= l	"Single nucleotide polymorphism"
CC	FT	variation	/standard_name=	replace(226,G)
CC	FT	variation	/*tag= m	"Single nucleotide polymorphism"
CC	FT	variation	/standard_name=	replace(230,C)
CC	FT	variation	/*tag= n	"Single nucleotide polymorphism"
CC	FT	variation	/standard_name=	replace(234,C)
XX	FT	variation	/*tag= o	"Single nucleotide polymorphism"
XX	FT	variation	/standard_name=	replace(236,C)
XX	FT	variation	/*tag= p	"Single nucleotide polymorphism"
XX	FT	variation	/standard_name=	replace(249,G)
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XX	FT	variation	/standard_name=	replace(281,C)
XX	FT	variation	/*tag= r	"Single nucleotide polymorphism"
XX	FT	variation	/standard_name=	replace(284,A)
XX	FT	variation	/*tag= s	"Single nucleotide polymorphism"
XX	FT	variation	/standard_name=	replace(284,C)
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XX	FT	variation	/standard_name=	replace(286,C)
XX	FT	variation	/*tag= u	"Single nucleotide polymorphism"
XX	FT	variation	/standard_name=	replace(303,C)
XX	FT	variation	/*tag= v	"Single nucleotide polymorphism"
XX	FT	variation	/standard_name=	replace(313,A)
XX	FT	variation	/*tag= w	"Single nucleotide polymorphism"
XX	FT	variation	/standard_name=	replace(508)
XX	FT	variation	/*tag= x	"Deletion of base A"
XX	FT	variation	/note=	replace(519,T)
XX	FT	variation	/*tag= y	"Single nucleotide polymorphism"
XX	FT	variation	/standard_name=	replace(524,A)
XX	FT	variation	/*tag= z	"Single nucleotide polymorphism"
XX	FT	variation	/standard_name=	replace(558,G)
XX	FT	variation	/*tag= aa	"Single nucleotide polymorphism"
XX	FT	variation	/standard_name=	replace(565,G)

Detecting growth hormone variants (GHI), useful in screening patients for growth hormone irregularities, comprises comparing the nucleotide sequence of a GHI gene from a test sample with that of a standard sequence of the human GHI

Example 2; Page 39; 95pp; English.

The invention described a method of detecting variation in growth hormone 1 (GHI), and therefore GH dysfunction in an individual. The method comprises comparing the nucleotide sequence of GHI gene obtained from the test sample with a standard human GHI gene sequence, in order to identify variation (GHI variant). The method is useful in screening patients for growth hormone irregularities or producing variant proteins for treating irregularities, and for the early detection and appropriate clinical management of familial GH deficiency. The GHI variants are useful in therapeutic, diagnostic or detection method, particularly for determining binding defects and susceptibility to a disease such as diabetes, obesity or infection; for treating acromegaly or gigantism conditions associated with lactogenic, diabetogenic, lipolytic and protein anabolic effects, syndromes, mood and sleep disorders; diagnosing GH dysfunction and determining pituitary storage defects. The GHI variants are especially useful in gene therapy or protein therapy. The GHI or GH variant may also be used in the preparation of a medicament, diagnostics composition or kit, or detection kit. The method has the advantage of: expanding the know spectrum of GHI gene mutations; evaluating the role of GHI gene mutations in the etiology of short stature; identifying of the mode of inheritance of novel lesions; evaluation the effects of GHI mutations on the structure and function of the GH molecule and development of rapid diagnostic tests for inherited GH deficiency. This sequence is the GHI PCR primer, GHPR, used with GHF (AAS18882) to amplify a GHI-specific fragment, described in the method of the invention.

Sequence 19 BP; 3 A; 2 C; 9 G; 5 T; 0 other;

Query Match 100.0%; Score 19; DB 24; Length 19;

Best Local Similarity 100.0%; Pred. No. 4.9;

Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 TGTAGGAAGTCTGGGTGC 19  
|||||  
DB 1 TGTAGGAAGTCTGGGTGC 19

RESULT 2

AAS18886/C

ID AAS18886 standard; DNA: 3700 BP.

AC AAS18886;

XX 12-MAR-2002 (first entry)

XX Growth hormone 1 gene (GHI), reference sequence.

DE Growth hormone 1; GHI; osteopathic; gene therapy; protein therapy;

XX Diabetes; obesity; infection; acromegaly; gigantism; sodium retention;

KW water retention; metabolic syndrome; mood disorder; sleep disorder;

KW Growth hormone dysfunction; familial growth hormone deficiency;

KW short stature; pituitary storage defect; human; chromosome 17q23; ds.

XX Homo sapiens.

OS

XX Key Location/Qualifiers

FT	FT	seq\_peptide	763..1100	
FT	FT	/\*tag= a		
FT	FT	CDS	763..2230	
FT	FT	/\*tag= b		
FT	FT	/product= "GHI"		
FT	FT	/note= "Growth hormone 1"		
FT	FT	variation	replace(124,G)	
FT	FT	/\*tag= c		
FT	FT	/standard\_name=	"Single nucleotide polymorphism"	

FT	variation	/standard_name= replace(1169,A) /*tag= ba	"Single nucleotide polymorphism"
FT		/standard_name= /*tag= ba	"Single nucleotide polymorphism"
FT	variation	replace(1182,T) /*tag= bb	"Single nucleotide polymorphism"
FT		/standard_name= replace(1189,G) /*tag= bc	"Single nucleotide polymorphism"
FT	variation	replace(1193,G) /*tag= bd	"Single nucleotide polymorphism"
FT	intron	/standard_name= 1194..1402 /*tag= be	"Single nucleotide polymorphism"
FT		/number= 2 replace(1196,G) /*tag= bf	"Single nucleotide polymorphism"
FT	variation	/standard_name= replace(1196,C) /*tag= bg	"Single nucleotide polymorphism"
FT		/standard_name= replace(1208,C) /*tag= bh	"Single nucleotide polymorphism"
FT	variation	/standard_name= replace(1212,T) /*tag= bi	"Single nucleotide polymorphism"
FT		/standard_name= /*tag= bi	"Single nucleotide polymorphism"

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Query Match      100.0%; Score 19; DB 24; length 3700;
Best local Similarity 100.0%; Pred. No. 6,4;
Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy      1 TGTAGGAAGTCTGGGGTGC 19
          |||||
Db      3316 TGTAGGAAGTCTGGGGTGC 3298

RESULT 3
AAS46366
ID      AAS46366 standard; DNA; 5333 BP.
XX
AC      AAS46366;
XX
DT      18-DEC-2001 (first entry)
XX
DE      Tumour suppressor gene derived chemically modified sequence #88.
XX
KW      Human; Tumour suppressor gene; oncogene; antitumour; cytostatic;
KW      cancer; tumour; CpG dinucleotide; single-nucleotide polymorphism; SNP;
KW      cytosine methylation; ds.
XX
OS      Homo sapiens.
XX
PN      W0200168912-A2.
XX
PD      20-SEP-2001.
XX
PF      15-MAR-2001; 2001W0-EP02955.
XX
PR      15-MAR-2000; 2000DE-1013847.
PR      06-APR-2000; 2000DE-1019058.
PR      07-APR-2000; 2000DE-1019173.
PR      30-JUN-2000; 2000DE-1032529.
PR      01-SEP-2000; 2000DE-1043826.
XX
PA      (EPiG-) EPiGENOMICS AG.
XX
PI      Olek A, Piepenbrock C, Berlin K;
XX
DR      WPI: 2001-602752/68.
XX
PT      Fragments of chemically modified genes associated with tumour suppressor

```

XX (EP)(G-) EPIGENOMICS AG.  
XX PPA  
XX Olek A. Piepenbrock C, Berlin K;  
XX PPI  
XX WPI; 2002-130909/17.  
XX  
XX Nucleic acid comprising fragment of chemically modified gene, useful  
XX for diagnosis and treatment of diseases associated with abnormal  
PT cytosine methylation -  
PT  
XX Claim 1; SEQ ID NO 756; 32pp + Sequence Listing; German.  
XX  
XX The present invention provides a number of human immune system associated  
CC genes which are modified by the methylation of cytosines. The sequences  
CC can be used in the diagnosis and treatment of immune system disorders,  
CC including eye diseases such as retinopathy, neovascular glaucoma and  
CC macular degeneration, arteriosclerosis, anaemia, cancer, acute myeloid  
CC leukaemia, Alzheimer's disease, AIDS, epilepsy, neurofibromatosis,  
CC rheumatoid arthritis, psoriasis and inflammatory/ulcerative bowel  
CC diseases. The present sequence is a gene of the invention.  
XX  
XX sequence 5333 BP; 1275 A; 101 C; 1345 G; 2612 T; 0 other;  
XX

QY 1 TGTAGGAAGTCCTGGGGTG 18  
|||||  
DB 4336 TGTAGGAAGTCCTGGGGTG 4353

RESULT 5  
ABN31023/c  
ID ABN31023 standard; DNA: 65 BP.

15-JUL-2002 (first entry)

Human; mouse; rat; splice transcript; detection; RNA transcript; splice variant; transcriptome; oligonucleotide library; ss.

[illegible]

07-FEB-2002

XX  
DE 30-1111-2001-2001WO-TB01903-[illegible]

PR 02-MAY-2001; 2001US-287724P.

PA (COMP-) COMPUGEN INC.

Shoshan A., Wasserman A., Mintz E., Mintz I., Faigler S;

WPT: 2002-257383/30

New oligonucleotide libraries comprising oligonucleotides which selectively hybridize to mRNAs transcribed from a transcription unit of a genome, useful for detecting tissue-, pathology-, and developmental-specific genes -

XX  
PS  
Example 1: SEQ ID 3771: 47bp: English.

transcription units that populate a genome. The library comprises several oligonucleotides, each capable of hybridizing selectively to a set of messenger RNAs transcribed from a given transcription unit of the genome, which encodes one or more messenger RNA splice variants. The oligonucleotide libraries are useful for detecting mRNAs from a biological sample, in expression profiling studies, in qualitatively or quantitatively characterising the corresponding transcriptome, and in detecting RNA transcripts and splice variants of human or animal transcriptomes. The libraries may also be used as specialised mini libraries to detect transcripts of a sub-transcriptome under a particular biological or pathological state, and so allowing the detection of tissue- and pathology-specific genes such as those genes only expressed in specific tissue under a specific pathological condition; to detect developmental specific genes; and to detect RNA transcripts and splice variants of a transcriptome of a patient suffering from a particular disorder. ABN27253 to ABN59589 represent oligonucleotide sequences from rats, humans and mice, which are used in the exemplification of the present invention.

N.B. The sequence data for this patent did not form part of the printed specification, but was obtained in electronic format directly from WIPO at [ftp.wipo.int/pub/published\\_pct\\_sequences](http://ftp.wipo.int/pub/published_pct_sequences).

Sequence 65 BP; 19 A; 12 C; 21 G; 13 T; 0 other;

Query Match 83.2%; Score 15.8; DB 24; Length 65;  
Best Local Similarity 89.5%; Pred. No. 2e+02;  
Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Oy 1 TGTAGGAAGTCTGGGTGC 19  
|||||  
Db 53 TGTAGGAAGTCTGGGTGC 35

RESULT 6  
AAK54271/c  
ID AAK54271 standard; cDNA; 428 BP.

XX AC AAK54271;

XX DT 16-NOV-2001 (first entry)

XX DE Murine transport and binding associated protein encoding cDNA SEQ ID 836.

XX KW Murine; liver; gene library; amino acid synthesis; binding protein;  
cell metabolism; energy metabolism; fatty acid metabolism; synthesis;  
phospholipid metabolism; purine; pyrimidine; nucleoside; nucleotide;  
replication; transcription; translation; transport protein; ss.

XX OS Mus musculus.

XX PN DE20103510-U1.

XX PD 07-JUN-2001.

XX PF 28-FEB-2001; 2001DE-2003510.

XX PR 02-DEC-1999; 99DE-1058160.

XX PA (LION-) LION BIOSCIENCE AG.

XX WPI; 2001-366570/39.

XX Gene library containing sequences with specific 3'-ends and no polyA tail, encoding proteins involved in a wide range of cellular processes

XX Claim 15; Page 250; 251pp; German.

XX This invention describes a novel gene library (A) comprises a gene sequence (or its part) encoding a protein involved in amino acid synthesis, cellular/energy metabolism, metabolism of fatty acids/phospholipids, synthesis or breakdown of purines/pyrimidines/nucleosides/nucleotides, DNA

replication/transcription/translation, or is a transport/binding protein. (A) are produced that correspond to the 3'-end of mRNA but without the polyA tail. They can be prepared more efficiently and with less effort than conventional libraries. AAK53436-AAK54275 represent fragments of the gene library described in the method of the invention.

Sequence 428 BP; 108 A; 99 C; 122 G; 99 T; 0 other;

Query Match 83.2%; Score 15.8; DB 22; Length 428;  
Best Local Similarity 89.5%; Pred. No. 2.2e+02;  
Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Oy 1 TGTAGGAAGTCTGGGTGC 19  
|||||  
Db 126 TGAAGGAAGTCTGGGTGC 108

RESULT 7  
AAF08345  
ID AAF08345 standard; cDNA; 730 BP.

XX AC AAF08345;

XX DT 13-MAR-2001 (first entry)

XX DE Fusarium venenatum EST SEQ ID NO: 868.

XX KW Multiple gene expression; filamentous fungal cell; EST;  
expressed sequence tag; Fusarium venenatum; Aspergillus niger;  
Aspergillus oryzae; Trichoderma reesei; identification; recombination;  
culture condition; environmental stress; spore morphogenesis;  
metabolic pathway engineering; catabolic pathway engineering; ss.

XX OS Fusarium venenatum.

XX PN WO2000056762-A2.

XX PD 28-SEP-2000.

XX PF 22-MAR-2000; 2000WO-US07781.

XX PR 22-MAR-1999; 99US-0273623.

XX PA (NOVO ) NOVO NORDISK BIOTECH INC.  
(NOVO ) NOVO NORDISK AS.

XX PI Berka RM, Key MW, Shuster JR, Kauppinen S, Clausen IG, Olsen PB;

XX WPI; 2000-594572/56.

XX Monitoring differential expression of genes in filamentous fungal cells uses fluorescence-labeled nucleic acids isolated from the cells and a substrate of expressed sequence tags.

XX Claim 86; Page 715; 3161pp; English.

XX The present invention describes a method for monitoring differential expression of genes in a first filamentous fungal (FF) cell relative to expression of the same genes in one or more second filamentous fungal cells. The method uses fluorescence-labeled nucleic acids isolated from the FF cells and a substrate of expressed sequence tags (EST). The ESTs are used in the methods for monitoring differential expression of genes in a first filamentous fungal (FF) cell relative to expression of the same genes in one or more second filamentous fungal cells. Monitoring the global expression of genes from FF cells allows the production potential of the microorganisms to be improved. New genes may be discovered, possible functions of unknown open reading frames can be identified and gene copy number variation and stability can be monitored. The expression of genes can be used to study how FF cells adapt to changes in culture conditions, environmental stress, spore morphogenesis, recombination, metabolic or catabolic pathway engineering. Using ESTs provides several advantages over genomic or random cDNA clones including elimination of redundancy as one spot on an



CC different from the wild type BMP1B polypeptide sequence and which has  
CC the ability to modulate ovulation in a female mammal. Mutation in  
CC the BMP1B receptor gene is responsible for increased ovulation rate  
CC in sheep derived from Booroola Merino strain that carry an autosomal  
CC mutation in FecR/Booroola gene. The FecB gene is mapped to chromosome 6.  
CC The BMP1A receptor of the invention and the polynucleotide encoding  
CC it are useful for modulating the ovulation rate of a female vertebrate.  
CC Identification of mutated BMP1B receptor nucleic acid molecule in a  
CC vertebrate, is useful for assessing fecundity in vertebrate such as  
CC humans and other commercially important mammals and birds including  
CC sheep, cattle, horses, goats, deer, pigs, cats, dogs, possums, and  
CC poultry. The polypeptide is useful to raise antibodies and for reducing  
CC unwanted populations of feral vertebrates. The polynucleotide is useful  
CC for identifying sequence variants in individual animals that are  
CC associated with increased ovulation. The present sequence is mutated  
CC BMP1B receptor cDNA from Booroola sheep. The BMP1B receptor is a  
CC member of transforming growth factor-beta family.  
XX  
SQ Sequence 1612 BP; 472 A; 359 C; 395 G; 386 T; 0 other;  
  
Query Match 83.2%; Score 15.8; DB 22; Length 1612;  
Best Local Similarity 89.5%; Pred. No. 2.3e+02;  
Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;  
  
QY 1 TGTAGGAAGTCTGGGGTGC 19  
DB 420 TGGAGGAAGTCTGGGGTGC 402  
IIIIIIIIIIIIIIIIIIII  
  
RESULT 10  
AA074064/C  
ID AA074064 standard; cDNA to mRNA; 2246 BP.  
XX  
AC AA074064;  
XX  
XX  
DT 29-JAN-1996 (first entry)  
DE Rat serotonin-lc cDNA.  
XX Serotonin; primer; mRNA; specificity; pharmaceutical; ss.  
KW Rattus rattus.  
OS JP07123984-A.  
PN 16-MAY-1995.  
PD 05-NOV-1993; 93JP-0275852.  
PF 05-NOV-1993; 93JP-0275852.  
XX (HITB ) HITACHI CHEM CO LTD.  
PA WPI; 1995-211627/28.  
DR  
XX A primer for the detection and the determin. of a specific messenger  
PT RNA - can detect and determine specific mRNA(s) with high  
PT reliability  
XX  
PS Example 28; Page 30-31; 35pp; Japanese.  
XX  
XX AA074064 is rat serotonin lc cDNA. This cDNA is amplified by the  
CC primers AA074047 and AA074048. The primers are used specifically for the  
CC detection and isolation of this sequence. They have the advantage of  
CC high sensitivity and reliability and are useful in the pharmaceutical  
CC industry.  
XX  
SQ Sequence 2246 BP; 532 A; 542 C; 544 G; 628 T; 0 other;  
  
Query Match 83.2%; Score 15.8; DB 16; Length 2246;  
Best Local Similarity 89.5%; Pred. No. 2.4e+02;  
Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 TGTAGGAAGTCTGGGGTGC 19  
DB 2096 TGTAGGAAGTCTGGGGTGC 2078  
IIIIIIIIIIIIIIIIIIII  
  
RESULT 11  
AAV48290/C  
ID AAV48290 standard; DNA; 2246 BP.  
XX  
AC AAV48290;  
XX  
DT 21-JAN-1999 (first entry)  
DE Rat 5-HT2C serotonin receptor S312F mutant DNA.  
XX  
KW Rat 5-HT2C serotonin receptor; G-protein-coupled receptor;  
KW constitutively activated monoamine G protein-coupled receptor;  
KW screen; agonist; inverse agonist; antagonist; mutant; ss.  
XX Synthetic.  
OS Rattus sp.  
XX  
FH Key Location/Qualifiers  
FT CDS 688..2070  
FT /\*Lag= a  
FT /product= Mutant 5-HT2C serotonin receptor  
FT 1621..1623  
FT /\*Lag= b  
FT /note= "codon has been changed from TCC to TTC"  
XX  
XX WO9838217-A1.  
XX  
XX 03-SFP-1998.  
XX 27-FEB-1998; 98WO-US03991.  
XX 07-OCT-1997; 97US-0061268.  
PR 27-FEB-1997; 97US-0039465.  
XX  
XX (EGAN/) EGAN C C.  
PA (HERR/) HERRICK-DAVIS K.  
PA (TEIT/) TEITLER M.  
XX Egan CC, Herrick-Davis K, Teitler M;  
XX  
XX WPI; 1998-495389/42.  
DR P-PSDB; AAW77113.  
XX  
XX Method of constitutively activating targeted G-protein coupled  
PT monoamine receptor - comprises use of site directed mutagenesis,  
PT useful for, e.g. screening for agonists and antagonists of native  
PT receptor  
XX  
XX Example 2; Fig 35; 97pp; English.  
XX  
XX The present sequence encodes a rat 5-HT2C serotonin receptor S312F  
CC mutant. 5-HT2C is a G-protein-coupled receptor. The sequence of the  
CC wild type receptor (AAV48279) is modified using the method of the  
CC invention to produce the present sequence. The mutant receptor exists  
CC in a constitutively activated state exhibiting both a greater response  
CC to agonists and a coupling to the G protein second messenger system  
CC even in the absence of agonists. The constitutively activated monoamine  
CC G protein-coupled receptor can be used to screen for agonists, inverse  
CC agonists, and antagonists of the native receptor.  
XX  
SQ Sequence 2246 BP; 537 A; 542 C; 538 G; 629 T; 0 other;  
  
Query Match 83.2%; Score 15.8; DB 19; Length 2246;  
Best Local Similarity 89.5%; Pred. No. 2.4e+02;  
Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;  
  
QY 1 TGTAGGAAGTCTGGGGTGC 19  
IIIIIIIIIIIIIIIIIIII

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Db 2096 TGTAGGAAGTCTGGCTGC 2078
RESULT 12
AAV48291/c
ID AAV48291 standard; DNA: 2246 BP.
XX AC AAV48291;
XX DT 21-JAN-1999 (first entry)
XX DE Rat 5-HT2C S312F mutant receptor DNA with a unique restriction site.
XX KW Rat 5-HT2C serotonin receptor; G-protein-coupled receptor;
XX KW constitutively activated monamine G protein-coupled receptor;
XX KW screen; agonist; inverse agonist; antagonist; mutant; ss.
XX OS Synthetic.
XX OS Rattus sp.
XX FH Key Location/Qualifiers
XX CDS 688..2070
XX FT /*tag= a
XX FT /product= Mutant 5-HT2C serotonin receptor
XX FT 1621..1623
XX FT /*tag= b
XX FT /note= "codon has been changed from TCC to TTC"
XX FT mutation
XX FT 1629
XX FT /*tag= c
XX FT /note= "C changed to A to create a ScaI site"
XX PN W09838217-A1.
XX PD 03-SEP-1998.
XX PF 27-FEB-1998; 98WO-US03991.
XX PR 07-OCT-1997; 97US-0061268.
XX PR 27-FEB-1997; 97US-0039465.
XX PA (EGAN/) EGAN C. C.
XX PA (HERR/) HERRICK-DAVIS K.
XX PA (TEIT/) TEITLER M.
XX PI Egan CC, Herrick-Davis K, Teitler M;
XX DR WPI: 1998-495389/42.
XX PT Method of constitutively activating targeted G-protein coupled
XX PT monoamine receptor - comprises use of site directed mutagenesis,
XX PT useful for, e.g. screening for agonists and antagonists of native
XX PT receptor
XX PS Claim 20; Fig 36; 97pp; English.
XX CC The present sequence encodes a rat 5-HT2C serotonin receptor S312F
XX CC mutant. The present sequence contains a ScaI restriction site. 5-HT2C
XX CC is a G-protein-coupled receptor. The sequence of the wild type receptor
XX CC (AAV48279) is modified using the method of the invention to produce the
XX CC present sequence. The mutant receptor exists in a constitutively
XX CC activated state exhibiting both a greater response to agonists and a
XX CC coupling to the G protein second messenger system even in the absence
XX CC of adonists. The constitutively activated monamine G protein-coupled
XX CC receptor can be used to screen for agonists, inverse agonists, and
XX CC antagonists of the native receptor.
XX SO Sequence 2246 BP; 538 A; 541 C; 538 G; 629 T; 0 other;
Query Match 83.2%; Score 15.8; DB 19; Length 2246;
Best Local Similarity 89.5%; Pred. No. 2.4e+02;
Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
Oy 1 TGTAGGAAGTCTGGCTGC 19
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Db 2096 TGTAGGAAGTCTGGCTGC 2078
RESULT 13
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ID AAV48288 standard; DNA: 2246 BP.
XX AC AAV48288;
XX DT 21-JAN-1999 (first entry)
XX DE Rat 5-HT2C serotonin receptor S312K mutant DNA.
XX KW Rat 5-HT2C serotonin receptor; G-protein-coupled receptor;
XX KW constitutively activated monamine G protein-coupled receptor;
XX KW screen; agonist; inverse agonist; antagonist; mutant; ss.
XX OS Synthetic.
XX OS Rattus sp.
XX FH Key Location/Qualifiers
XX CDS 688..2070
XX FT /*tag= a
XX FT /product= Mutant 5-HT2C serotonin receptor
XX FT 1621..1623
XX FT /*tag= b
XX FT /note= "codon has been changed from TCC to AAG"
XX PN W09838217-A1.
XX PD 03-SEP-1998.
XX PF 27-FEB-1998; 98WO-US03991.
XX PR 07-OCT-1997; 97US-0061268.
XX PR 27-FEB-1997; 97US-0039465.
XX PA (EGAN/) EGAN C. C.
XX PA (HERR/) HERRICK-DAVIS K.
XX PA (TEIT/) TEITLER M.
XX PI Egan CC, Herrick-Davis K, Teitler M;
XX DR WPI: 1998-495389/42.
XX DR P-PSDB; AAW77112.
XX PT Method of constitutively activating targeted G-protein coupled
XX PT monoamine receptor - comprises use of site directed mutagenesis,
XX PT useful for, e.g. screening for agonists and antagonists of native
XX PT receptor
XX PS Example 2; Fig 32; 97pp; English.
XX CC The present sequence encodes a rat 5-HT2C serotonin receptor S312K
XX CC mutant. 5-HT2C is a G-protein-coupled receptor. The sequence of the
XX CC wild type receptor (AAV48279) is modified using the method of the
XX CC invention to produce the present sequence. The mutant receptor exists
XX CC in a constitutively activated state exhibiting both a greater response
XX CC to agonists and a coupling to the G protein second messenger system
XX CC even in the absence of agonists. The constitutively activated monamine
XX CC G protein-coupled receptor can be used to screen for agonists, inverse
XX CC agonists, and antagonists of the native receptor.
XX SO Sequence 2246 BP; 539 A; 541 C; 539 G; 627 T; 0 other;
Query Match 83.2%; Score 15.8; DB 19; Length 2246;
Best Local Similarity 89.5%; Pred. No. 2.4e+02;
Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
Oy 1 TGTAGGAAGTCTGGCTGC 19
Db 2096 TGTAGGAAGTCTGGCTGC 2078
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Thu Jan 23 13:47:06 2003

us-09-853-688-36.rng

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RESULT 14
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ID AAV48289 standard; DNA: 2246 BP.
XX
XX AAV48289;
XX
DT 21-JAN-1999 (first entry)
XX
XX Rat 5-HT2C S312K mutant receptor DNA with a unique restriction site.
DE
XX
XX Rat 5-HT2C serotonin receptor; G-protein-coupled receptor;
KW constitutively activated monoamine G protein-coupled receptor;
KW screen; agonist; inverse agonist; antagonist; mutant; ss.
XX
XX Synthetic.
OS Rattus sp.
XX
XX Key Location/Qualifiers
FH 688..2070
FT CDS
FT /*tag= a
FT /product= Mutant 5-HT2C serotonin receptor
FT 1621..1623
FT /*tag= b
FT /note= codon has been changed from TCC to AAG"
FT mutation
FT 1629
FT /*tag= c
FT /note= *C changed to A to create a ScaI site"
FT mutation
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XX WO9838217-A1.
PN
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XX 03-SEP-1998.
PD
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XX 27-FEB-1998; 98WO-US039465.
PF
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XX 07-OCT-1997; 97US-0061268.
PR
XX 27-FEB-1997; 97US-0039465.
PR
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XX (EGAN/) EGAN C C.
PA (HERR/) HERRICK-DAVIS K.
PA (TEIT/) TEITLER M.
XX
XX Egan CC, Herrick-Davis K, Teitler M;
PI
XX WPI; 1998-495389/42.
XX
XX Method of constitutively activating targeted G-protein coupled
PT monoamine receptor - comprises use of site directed mutagenesis,
PT useful for, e.g. screening for agonists and antagonists of native
PT receptor
XX
XX Claim 19; Fig 33; 97pp; English.
XX
XX The present sequence encodes a rat 5-HT2C serotonin receptor S312K
CC mutant. The present sequence contains a ScaI restriction site. 5-HT2C
CC is a G-protein-coupled receptor. The sequence of the wild type receptor
CC (AAV48279) is modified using the method of the invention to produce the
CC present sequence. The mutant receptor exists in a constitutively
CC activated state exhibiting both a greater response to agonists and a
CC coupling to the G protein second messenger system even in the absence
CC of agonists. The constitutively activated monoamine G protein-coupled
CC receptor can be used to screen for agonists, inverse agonists, and
CC antagonists of the native receptor.
XX
XX Sequence 2246 BP; 540 A; 540 C; 539 G; 627 T; 0 other;
SQ
Query Match 83.2%; Score 15.8; DB 19; Length 2246;
Best Local Similarity 89.5%; Pred. NO. 2.4e+02;
Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
OY 1 TGTAGTAAGTCTGGGGTGC 19
|||||

```

```
Db 2096 TGTAGGAAGTCTGCGCTGC 2078
RESULT 15
AAV48279/c
ID AAV48279 standard; DNA: 2246 BP.
XX
XX AAV48279;
XX
DT 21-JAN-1999 (first entry)
XX
XX Rat 5-HT2C serotonin receptor DNA.
DE
XX
XX Rat 5-HT2C serotonin receptor; G-protein-coupled receptor;
KW constitutively activated monoamine G protein-coupled receptor;
KW screen; agonist; inverse agonist; antagonist; ss.
XX
XX Synthetic.
OS Rattus sp.
XX
XX Key Location/Qualifiers
FH 688..2070
FT CDS
FT /*tag= a
FT /product= 5-HT2C serotonin receptor
FT 1621..1623
FT /*tag= b
FT /note= codon has been changed from TCC to AAG"
FT mutation
FT 1629
FT /*tag= c
FT /note= *C changed to A to create a ScaI site"
FT mutation
XX
XX WO9838217-A1.
PN
XX
XX 03-SEP-1998.
PD
XX
XX 27-FEB-1998; 98WO-US039991.
PF
XX
XX 07-OCT-1997; 97US-0061268.
PR
XX 27-FEB-1997; 97US-0039465.
PR
XX
XX (EGAN/) EGAN C C.
PA (HERR/) HERRICK-DAVIS K.
PA (TEIT/) TEITLER M.
XX
XX Egan CC, Herrick-Davis K, Teitler M;
PI
XX WPI; 1998-495389/42.
XX
XX Method of constitutively activating targeted G-protein coupled
PT monoamine receptor - comprises use of site directed mutagenesis,
PT useful for, e.g. screening for agonists and antagonists of native
PT receptor
XX
XX Disclosure; Fig 2A; 97pp; English.
XX
XX The present sequence encodes a rat 5-HT2C serotonin receptor. This
CC is a G-protein-coupled receptor. The sequence of the receptor is
CC modified (see AAV48288-91) using the method of the invention so
CC that the receptor exists in a constitutively activated state exhibiting
CC both a greater response to agonists and a coupling to the G protein
CC second messenger system even in the absence of agonists. The
CC constitutively activated monoamine G protein-coupled receptor can be
CC used to screen for agonists, inverse agonists, and antagonists of the
CC native receptor.
XX
XX Sequence 2246 BP; 537 A; 543 C; 538 G; 628 T; 0 other;
SQ
Query Match 83.2%; Score 15.8; DB 19; Length 2246;
Best Local Similarity 89.5%; Pred. NO. 2.4e+02;
Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
OY 1 TGTAGGAAGTCTGCGGTGC 19
|||||
Db 2096 TGTAGGAAGTCTGCGCTGC 2078
Search completed: January 21, 2003, 16:28:00
Job time : 161.389 secs

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GenCore version 5.1.3  
Copyright (c) 1993 - 2003 Compuen Ltd.

(M) nucleic - nucleic search, using sw model

Run on: January 21, 2003, 14:52:32 : Search time 1198.97 Seconds  
(without alignments)  
229.633 Million cell updates/sec

Title: US-09-853-688-35

Perfect score: 17  
Sequence: 1 gggagccccagcaatgc 17

Scoring table: IDENTITY\_NUC  
Gapop 10.0 , Gapext 1.0

Searched: 16154066 seqs, 8097743376 residues

Total number of hits satisfying chosen parameters: 32308132

Minimum dB seq length: 0

Maximum DB seq length: 20000000000

Post-processing: Minimum Match 0%

Maximum Match 100%  
Listing first 45 summaries

Database :

EST:\*

- 1: em\_estba:\*
- 2: em\_esthum:\*
- 3: em\_estin:\*
- 4: em\_estov:\*
- 5: em\_estov:\*
- 6: em\_estpl:\*
- 7: em\_estro:\*
- 8: em\_hic:\*
- 9: gb\_estl:\*
- 10: gb\_est2:\*
- 11: gb\_hic:\*
- 12: gb\_est3:\*
- 13: gb\_est4:\*
- 14: gb\_est5:\*
- 15: em\_estfun:\*
- 16: em\_estom:\*
- 17: gb\_gss:\*
- 18: em\_gss\_hum:\*
- 19: em\_gss\_inv:\*
- 20: em\_gss\_pln:\*
- 21: em\_gss\_vrt:\*
- 22: em\_gss\_fun:\*
- 23: em\_gss\_mam:\*
- 24: em\_gss\_mus:\*
- 25: em\_gss\_other:\*
- 26: em\_gss\_pro:\*
- 27: em\_gss\_rod:\*

pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result, being printed, and is derived by analysis of the total score distribution.

#### SUMMARIES

Result No.	Score	Match	Length	DB	ID	Description
1	16	94.1	461	10	AW558892	L0301D12-
2	16	94.1	619	17	BH108836	RPCI-24-2
3	16	94.1	621	12	BG324158	602423116
4	16	94.1	636	10	AW239325	xb39a04.y
5	16	94.1	642	12	BG076347	H3158F07-
6	16	94.1	774	12	BG704650	602688222

c	7	16	94.1	976	13	BM468789
c	8	15.4	90.6	300	9	AA852498
c	9	15.4	90.6	319	10	AW067967
c	10	15.4	90.6	320	9	A1449767
c	11	15.4	90.6	336	9	AA629118
c	12	15.4	90.6	382	17	BH842232
c	13	15.4	90.6	382	17	BH845093
c	14	15.4	90.6	383	13	BT005357
c	15	15.4	90.6	385	17	BT14181
c	16	15.4	90.6	393	9	AT1752118
c	17	15.4	90.6	406	12	BF406319
c	18	15.4	90.6	411	9	AA536528
c	19	15.4	90.6	411	12	BF856554
c	20	15.4	90.6	412	14	R45494
c	21	15.4	90.6	413	9	AT1751985
c	22	15.4	90.6	413	12	BF857617
c	23	15.4	90.6	416	13	BT239628
c	24	15.4	90.6	420	9	AT1751499
c	25	15.4	90.6	424	9	AT1751854
c	26	15.4	90.6	425	10	BB268875
c	27	15.4	90.6	440	9	AT1751606
c	28	15.4	90.6	454	9	AA391288
c	29	15.4	90.6	457	9	AT1750858
c	30	15.4	90.6	468	13	BT243443
c	31	15.4	90.6	501	14	BM727488
c	32	15.4	90.6	512	13	BT363494
c	33	15.4	90.6	512	13	BT403236
c	34	15.4	90.6	537	13	BT371847
c	35	15.4	90.6	540	9	AU144696
c	36	15.4	90.6	549	13	BT238029
c	37	15.4	90.6	562	13	BT486096
c	38	15.4	90.6	581	14	W22315
c	39	15.4	90.6	584	9	AA160661
c	40	15.4	90.6	587	9	AT520081
c	41	15.4	90.6	589	12	BG162438
c	42	15.4	90.6	591	14	BM931980
c	43	15.4	90.6	594	14	BM685020
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c	45	15.4	90.6	607	13	BT162042

#### ALIGNMENTS

RESULT 1  
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LOCUS L0301D12-3 NIA Mouse Newborn Ovary CDNA Library Mus musculus CDNA  
DEFINITION clone L0301D12 3', mRNA sequence.  
ACCESSION AW558892  
VERSION AW558892.1 GI:7204321  
KEYWORDS EST.  
SOURCE house mouse.  
ORGANISM Mus musculus  
Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.  
REFERENCE Tanaka,T.S., Jaradat,S.A., Lim,M.K., Kargul,G.J., Wang,X., Grahovac,M.J., Pantano,S., Sano,Y., Piao,Y., Nagaraja,R., Dol,H., Wood,W.H., III, Becker,K.G. and KO,M.S.H.  
TITLE Genome-wide expression profiling of mid-gestation placenta and embryo using a 15,000 mouse developmental CDNA microarray  
JOURNAL Proc. Natl. Acad. Sci. U.S.A. 97 (16), 9127-9132 (2000)  
MEDLINE 20381348  
COMMENT Contact: George J. Kargul  
Laboratory of Genetics  
National Institute on Aging/National Institutes of Health  
333 Cassell Drive, Suite 4000, Baltimore, MD 21224-6820, USA  
Email: cdna@lgsun.grc.nia.nih.gov  
Plate: L0301 row: D column: 12  
Seq primer: -21M3 Forward  
High quality sequence stop: 461  
POLYA=Yes.

FEATURES  
source

Location/Qualifiers  
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/strain="C57Bl/6J"  
/db\_xref="nlcst:L0301012-3"  
/db\_xref="taxon:10090"  
/clone="L0301012"  
/clone\_lib="NIA Mouse Newborn Ovary cDNA Library"  
/sex="female"  
/dev\_stage="Newborn Ovary"  
/lab\_host="DH10B"  
/note="Vector: pSPORT1 (Gibco/BRL Life Technology); Site\_1: SalI; Site\_2: NotI; Total RNAs were extracted from 7 Newborn Ovary. The double-stranded cDNA was synthesized by Gibco's kit with an Oligo(dT) primer [NotI primer-adaptor from GibcoBRL] 15'-GCAGTCTCTAGATCGAGCGCCGCTTTT-3' from 2.56ug of total RNA. The double-stranded cDNAs were treated with T4 DNA polymerase and purified by ethanol-precipitation. The cDNAs were ligated to lone-linker L1-SalI (include SalI sequence). The cDNAs were purified by phenol/chloroform and separated from free linkers by Centricon 100. Then, cDNAs were amplified by long-range high fidelity PCR using Takara's Ex Taq polymerase. Then, the cDNAs were purified by phenol/chloroform and by Centricon 100. The cDNAs were digested with SalI and NotI enzymes. Then, the cDNAs were size selected by Gibco's Size Fractionation Column. The cDNAs were cloned into SalI/NotI site of pSPORT1 plasmid vector. The DH10B E. coli host was transformed with the ligation mixture by chemical method. The library was constructed by Xiaohong Wang and Yulan Piao."

BASE COUNT 125 a 86 c 126 g 124 t

ORIGIN

Query Match 94.1%; Score 16; DB 10; Length 461;  
Best Local Similarity 100.0%; Pred. No. 2.1e+03;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GCGAGCCCCAGCAATG 16  
|||||

DB 240 GCGAGCCCCAGCAATG 295

RESULT 2  
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LOCUS  
DEFINITION  
ACCESSION  
VERSION  
KEYWORDS  
SOURCE  
ORGANISM

BH108836 619 bp DNA linear GSS 19-JUL-2001  
RPCI-24-238C2-TV RPCI-24 Mus musculus genomic clone RPCI-24-238C2,  
DNA sequence.  
BH108836 GI:14941302  
GSS.  
house mouse.  
Mus musculus  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.  
1 (bases 1 to 619)  
Zhao,S., Nierman,W., Malek,J., Shatsman,S., Akinret,B., Levins,M.,  
Tsegaye,G., Geer,K., Krol,M., Shvartsbeyn,A., Gebregeorgis,E.,  
Russell,D., de Jong,P. and Fraser,C.M.  
Mouse BAC End Sequences from Library RPCI-24  
Other\_GSSs: RPCI-24-238C2.TJ  
Contact: Shaying Zhao  
Department of Eukaryotic Genomics  
The Institute for Genomic Research  
9712 Medical Center Dr., Rockville, MD 20850, USA  
Tel: 301 838 0200  
Fax: 301 838 0208  
Email: szhao@tigr.org  
Clones are derived from the mouse BAC library RPCI-24. For BAC  
library availability, please contact Pieter de Jong  
(pdejong@mail.cho.org). Clones may be purchased from BACPAC

Resources (http://www.choi.org/bacpac/orderingframe.htm). BAC end  
page: http://www.tigr.org/tdb/bac\_ends/mouse/bac\_end\_intro.html  
Plate: 238 row: C column: 2  
Seq primer: T7  
Class: BAC ends.

FEATURES  
source

Location/Qualifiers  
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/strain="C57Bl/6J"  
/db\_xref="taxon:10090"  
/clone="RPCI-24-238C2"  
/clone\_lib="RPCI-24"  
/sex="Male"  
/cell\_type="Spleen/Brain"  
/note="Vector: pTAR8AC1; Site\_1: BamHI; Site\_2: BamHI;  
RPCI-24 Mouse BAC Library produced by Pieter de Jong. The  
library was cloned in the pTAR8AC1 cloning vector at the  
BamHI sites using MboI partially digested male C57Bl/6J  
DNA."

BASE COUNT 138 a 149 c 172 g 160 t

ORIGIN

Query Match 94.1%; Score 16; DB 17; Length 619;  
Best Local Similarity 100.0%; Pred. No. 2.3e+03;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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|||||

DB 207 GCGAGCCCCAGCAATG 222

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LOCUS  
DEFINITION  
ACCESSION  
VERSION  
KEYWORDS  
SOURCE  
ORGANISM

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602423116F1 NIH\_MGC\_14 Homo sapiens cDNA clone IMAGE:4561371 5',  
mRNA sequence.  
BG324158  
EST.  
BG324158.1 GI:13130595  
human.  
Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.  
1 (bases 1 to 621)  
NIH-MGC http://mgc.nci.nih.gov/.  
National Institutes of Health, Mammalian Gene Collection (MGC)  
Unpublished (1999)  
Contact: Robert Strausberg, Ph.D.  
Email: cgapbs@mail.nih.gov  
Tissue Procurement: DCTD/DTP  
cDNA Library Preparation: Ling Hong/Rubin Laboratory  
cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)  
DNA Sequencing by: Incyte Genomics, Inc.  
Clone distribution: MGC clone distribution information can be  
found through the I.M.A.G.E. Consortium/LLNL at:  
http://image.llnl.gov  
Plate: LLCM1271 row: o column: 04  
High quality sequence stop: 615.  
Location/Qualifiers  
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/db\_xref="taxon:9606"  
/clone="IMAGE:4561371"  
/clone\_lib="NIH\_MGC\_14"  
/tissue\_type="renal cell adenocarcinoma"  
/lab\_host="DH10B (phage-resistant)"  
/note="Organ: kidney; Vector: pOT87; Site\_1: XhoI; Site\_2:  
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adaptor: GGCACGAG(G). Size-selected >500bp for average  
insert size 1.8kb. Library constructed by Ling Hong in  
the laboratory of Gerald M. Rubin (University of  
California, Berkeley) using ZAP-cDNA synthesis kit

FEATURES  
source

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BASE COUNT      128 a 185 c 188 g 120 t
ORIGIN
      (Stratagene) and Superscript II RT (Life Technologies).
Query Match      94.1%; Score 16; DB 12; Length 621;
Best Local Similarity 100.0%; Pred. No. 2.3e+03;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY  2  GGAGCCCGCAGCAATGC 17
      |||||
Db   41  GGAGCCCGCAGCAATGC 16

RESULT 4
LOCUS      AW239325/c
DEFINITION xB39a04_Y1 NCI_CGAP_Lu31 Homo sapiens cDNA clone IMAGE:2578638 5'
           similar to gb:J03934 NAD(P)H DEHYDROGENASE (HUMAN);, mRNA sequence.
ACCESSION  AW239325
VERSION     AW239325.1 GI:6571715
KEYWORDS   EST.
SOURCE     human.
ORGANISM   Homo sapiens
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE  1 (bases 1 to 636)
            NCI-CCGAP http://www.ncbi.nlm.nih.gov/ncicgap.
            National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
            Tumor Gene Index
JOURNAL    Unpublished (1997)
COMMENT    Other ESTs: xB39a04.x1
            Contact: Robert Strausberg, Ph.D.
            Email: cyaphs-r@mail.nih.gov
            Tissue Procurement: ATCC cDNA Library Preparation: Life
            Technologies, Inc. cDNA Library Arrayed by: Christa Prange, The
            I.M.A.G.E. Consortium DNA Sequencing by: Washington University
            Genome Sequencing Center
            Clone distribution: NCI-CCGAP clone distribution information can be
            found through the I.M.A.G.E. Consortium/LLNL at:
            www.bio.llnl.gov/bbrp/image/image.html
            Seq primer: -40RP from Gibco
            High quality sequence stop: 437.
FEATURES
    source
        1..636
        /organism="Homo sapiens"
        /db_xref="taxon:9606"
        /clone="IMAGE:2578638"
        /clone_lib="NCI_CGAP_Lu31"
        /sex="male"
        /dev_stage="fetal, 14 wk post-conception"
        /lab_host="DH10B"
        /note="Organ: lung, cell line; Vector: pCMV-Sport6;
        Site_1: EcorV; Site_2: Nott; Cloned unidirectionally, no
        5' adaptor. Primer: Oligo dT. Full-length library
        constructed by Life Technologies."
BASE COUNT      154 a 170 c 168 g 143 t 1 others
ORIGIN
      (Stratagene) and Superscript II RT (Life Technologies).
Query Match      94.1%; Score 16; DB 10; Length 636;
Best Local Similarity 100.0%; Pred. No. 2.3e+03;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY  2  GGAGCCCGCAGCAATGC 17
      |||||
Db   635  GGAGCCCGCAGCAATGC 620

RESULT 5
LOCUS      BG076347
DEFINITION H3158F07-3 NIA Mouse 15K cDNA Clone Set Mus musculus cDNA clone
            H3158F07 3', mRNA sequence.
ACCESSION  BG076347

```

```

VERSION          BG076347.1 GI:12558916
KEYWORDS         EST.
SOURCE           house mouse.
ORGANISM         Mus musculus
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE        1 (bases 1 to 642)
AUTHORS          Karqui,G.J., Dudekula,D.B., Qian,Y., Lim,M.K., Jaradat,S.A., Tanaka
            ,T.S., Carter,M.G. and Ko,M.S.H.
JOURNAL          Unpublished (2001)
COMMENT          Verification and initial annotation of NIA mouse 15K cDNA clone set
            Other ESTs: H3158F07-5
            Contact: George J. Karqui
            Laboratory of Genetics
            National Institute on Aging/National Institutes of Health
            333 Cassell Drive, Suite 4000, Baltimore, MD 21224-6820, USA
            Email: cdna@igsun.grc.nia.nih.gov
            This clone set has been freely distributed to the community. Please
            visit http://lgsun.grc.nia.nih.gov/cDNA/15k.html for details.
            Plate: H3158 row: F column: 07
            Seq primer: -21M13 Forward
            High quality sequence stop: 642
            POLYA=Yes.
FEATURES
    source
        1..642
        /organism="Mus musculus"
        /strain="C57BL/6J"
        /db_xref="niaEST:H3158F07-3"
        /db_xref="taxon:10090"
        /clone="H3158F07"
        /clone_lib="NIA Mouse 15K cDNA Clone Set"
        /sex="Clones arrayed from a variety of cDNA libraries"
        /dev_stage="Clones arrayed from a variety of cDNA
        libraries"
        /lab_host="DH10B"
        /note="Vector: pSPORT1; Site_1: SalI; Site_2: NotI; This
        clone is among a rearranged set of 15,247 clones from 11
        embryo cDNA libraries (including preimplantation stage
        embryos from unfertilized egg to blastocyst, embryonic
        part of E7.5 embryos, extraembryonic part of E7.5 embryos
        , and E12.5 female mesonephros/gonad) and one newborn
        ovary cDNA library. Average insert size 1.5 kb. All
        source libraries are cloned unidirectionally with Oligo(dT
        )-Not primers. References include: (1) Genome-wide
        expression profiling of mid-gestation placenta and embryo
        using a 15,000 mouse developmental cDNA microarray, 2000,
        Proc. Natl. Acad. Sci. U S A, 97: 9127-9132; (2)
        Large-scale cDNA analysis reveals phased gene expression
        patterns during preimplantation mouse development, 2000,
        Development, 127: 1737-1749; (3) Genome-wide mapping of
        unselected transcripts from extraembryonic tissue of
        7.5-day mouse embryos reveals enrichment in the t-complex
        and under-representation on the X chromosome, 1998, Hum
        Mol Genet 7: 1967-1978."
BASE COUNT      166 a 135 c 181 g 160 t
ORIGIN
      (Stratagene) and Superscript II RT (Life Technologies).
Query Match      94.1%; Score 16; DB 12; Length 642;
Best Local Similarity 100.0%; Pred. No. 2.3e+03;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY  1  GGGAGCCCGCAGCAATG 16
      |||||
Db   280  GGGAGCCCGCAGCAATG 295

RESULT 6
LOCUS      BG704650/c
DEFINITION 602688222P1 NIH_MGC_95 Homo sapiens cDNA clone IMAGE:4820860 5',
            mRNA sequence.
ACCESSION  BG704650
VERSION     BG704650.1 GI:13978201

```

EST.  
SOURCE human.  
ORGANISM Homo sapiens  
REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
AUTHORS Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.  
TITLE NIH-MGC http://mgc.nci.nih.gov/.  
JOURNAL 1 (bases 1 to 774)  
COMMENT National Institutes of Health, Mammalian Gene Collection (MGC)  
Unpublished (1999)  
Contact: Robert Strausberg, Ph.D.  
Email: cgapbs@mail.nih.gov  
Tissue Procurement: Miklos Palkovits, M.D., Ph.D.  
CDNA Library Preparation: Michael J. Brownstein (NHGRI), Shiraki  
Toshiyuki and Piero Carninci (RIKEN)  
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)  
DNA Sequencing by: Incyte Genomics, Inc.  
Clone distribution: MGC clone distribution information can be  
found through the I.M.A.G.E. Consortium/LLNL at:  
http://image.llnl.gov  
Plate: L1AM10726 row: k column: 05  
High quality sequence stop: 700.  
Location/Qualifiers  
1..774  
/organism="Homo sapiens"  
/db\_xref="taxon:9606"  
/clone="IMAGE:4820860"  
/clone\_lib="NIH\_MGC\_95"  
/tissue\_type="hippocampus"  
/lab\_host="DH10B"  
/note="Organ: brain; Vector: pBluescriptR (modified  
pBluescript KS+); Site:1: BamHI; Site:2: SalI-XhoI (gtcag  
); Oligo-dT primed using primer 5'-TTTTTTTTTTTNN-3',  
size-selected for average insert size 2.5 kb and  
normalized to 50x. This is a primary library enriched  
for full-length clones and constructed using the  
Cap-trapper method (Carninci, in preparation). Library  
constructed by M. Brownstein (NIMH/NHGRI, National  
Institutes of Health). Note: this is a NIH\_MGC Library."  
BASE COUNT 171 a 216 c 233 g 154 t  
ORIGIN  
Query Match 94.1%; Score 16; DB 12; Length 774;  
Best Local Similarity 100.0%; Pred. No. 2.4e+03;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 2 GGAGCCCCCAGCAATGC 17  
Db 54 GGAGCCCCCAGCAATGC 39  
RESULT 7  
HM468789/c 976 bp mRNA linear EST 05-FEB-2002  
LOCUS AGENCOURT\_6481239 NIH\_MGC\_88 Homo sapiens cDNA clone IMAGE:5541510  
DEFINITION 5', mRNA sequence.  
ACCESSION BM468789  
VERSION BM468789.1 GI:18517831  
KEYWORDS EST.  
SOURCE human.  
ORGANISM Homo sapiens  
REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
AUTHORS Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.  
TITLE NIH-MGC http://mgc.nci.nih.gov/.  
JOURNAL 1 (bases 1 to 976)  
COMMENT National Institutes of Health, Mammalian Gene Collection (MGC)  
Unpublished (1999)  
Contact: Robert Strausberg, Ph.D.  
Email: cgapbs@mail.nih.gov  
Tissue Procurement: ATCC  
CDNA Library Preparation: Life Technologies, Inc.  
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)  
DNA Sequencing by: Agencourt Bioscience Corporation  
Clone distribution: MGC clone distribution information can be

found through the I.M.A.G.E. Consortium/LLNL at:  
http://image.llnl.gov  
Plate: L1AM12239 row: f column: 07  
High quality sequence stop: 642.  
Location/Qualifiers  
1..976  
/organism="Homo sapiens"  
/db\_xref="taxon:9606"  
/clone="IMAGE:5541510"  
/clone\_lib="NIH\_MGC\_88"  
/tissue\_type="duodenal adenocarcinoma, cell line"  
/lab\_host="DH10B (phage-resistant)"  
/note="Organ: small intestine; Vector: pCMV-SPORT6;  
Site:1: NotI; Site:2: SalI; Cloned unidirectionally;  
oligo-dT primed. Average insert size 1.767 kb. Library  
enriched for full-length clones and constructed by Life  
Technologies. Note: this is a NIH\_MGC Library."  
BASE COUNT 202 a 289 c 267 g 218 t  
ORIGIN  
Query Match 94.1%; Score 16; DB 13; Length 976;  
Best Local Similarity 100.0%; Pred. No. 2.6e+03;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 1 GGAGCCCCCAGCAATG 16  
Db 705 GGAGCCCCCAGCAATG 690  
RESULT 8  
AA852498 300 bp mRNA linear EST 20-JUN-2002  
LOCUS NHTBCae14a07f1 Normal Human Trabecular Bone Cells Homo sapiens cDNA  
DEFINITION Clone NHTBCae14a07, mRNA sequence.  
ACCESSION AA852498  
VERSION AA852498.1 GI:2941091  
KEYWORDS EST.  
SOURCE human.  
ORGANISM Homo sapiens  
REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
AUTHORS Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.  
TITLE Jia, L.B., Young, M.F., Touchman, J.W., Bouffard, G.G., Robey  
Beckstrom-Stenberg, S.M., Green, E.D., Powell, J.I., Yang, L.M., Robey  
P.G., Hotchkiss, R.N. and Francomano, C.A.  
JGAP: The Skeletal Genome Anatomy Project  
Unpublished (1997)  
Contact: Libin Jia  
Medical Genetics Branch  
National Human Genome Research Institute  
10/10C101, 9000 Rockville Pike, Bethesda, MD 20892-1267, USA  
Tel: 301-496-7157  
Fax: 301-496-7157  
Email: libin@helix.nih.gov  
Seq primer: M13 Forward.  
Location/Qualifiers  
1..300  
/organism="Homo sapiens"  
/db\_xref="taxon:9606"  
/clone="NHTBCae14a07"  
/clone\_lib="Normal Human Trabecular Bone Cells"  
/sex="Female"  
/tissue\_type="Bone"  
/cell\_type="Trabecular Bone Cells"  
/lab\_host="SURE"  
/note="Organ: Hip; Vector: pBluescript; Site:1: EcoRI;  
Library constructed by Dr. Marian Young and Dr. Pamela  
Gehron Robey (NIDCR)"  
BASE COUNT 64 a 102 c 99 g 34 t 1 others  
ORIGIN  
Query Match 90.6%; Score 15.4; DB 9; Length 300;  
Best Local Similarity 94.1%; Pred. No. 3.5e+03;

Matches 16: Conservative 0: Mismatch 1: Indels 0: Gaps 0:

Qy 1 GGGAGCCCCAGCAATGC 17  
 |||||  
 Db 167 GGGAGCCCCAGCAACGC 183

RESULT 9  
 AW067967/c  
 LOCUS  
 DEFINITION  
 clone2h11.x1 Normal Human Trabecular Bone Cells Homo sapiens cDNA  
 clone NHTBC\_cn22h11 random, mRNA sequence.

ACCESSION  
 AW067967  
 VERSION  
 AW067967.1 GI:6022965  
 KEYWORDS  
 EST.  
 SOURCE  
 human.  
 ORGANISM  
 Homo sapiens  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE  
 AUTHORS  
 1 (bases 1 to 319)  
 Jia,L.B., Young,M.F., Touchman,J.W., Bouffard,G.G.,  
 Beckstrom-Sternberg,S.M., Green,E.D., Powell,J.I., Yang,L.M., Robey  
 P.G., Hotchkiss,R.N. and Francomano,C.A.  
 SGAP: The Skeletal Genome Anatomy Project  
 Unpublished (1997)  
 Contact: Libin Jia  
 Medical Genetics Branch  
 National Human Genome Research Institute  
 10/10C101, 9000 Rockville Pike, Bethesda, MD 20892-1267, USA  
 Tel: 301-402-4877  
 Fax: 301-496-7157  
 Email: libin@helix.nih.gov  
 DNA Sequencing and analyses by National Institutes of Health  
 Intramural Sequencing Center (NISC).  
 Plate: 22 row: h column: 11  
 Seq primer: 21M13 forward primer (ABI).  
 Location/Qualifiers

FEATURES  
 source  
 1..319  
 /organism="Homo sapiens"  
 /db\_xref="taxon:9606"  
 /clone="NHTBC\_cn22h11"  
 /clone\_lib="Normal Human Trabecular Bone Cells"  
 /sex="Female"  
 /tissue\_type="Bone"  
 /cell\_type="Trabecular Bone Cells"  
 /lab\_host="SURE"  
 /note="Organ: Hip; Vector: pBluescript; Site\_1: EcoRI;  
 Library constructed by Dr. Marian Young and Dr. Pamela  
 Gehron Robey (NIDCR)."  
 41 a 96 c 112 g 70 t

BASE COUNT  
 ORIGIN  
 Query Match 90.6%; Score 15.4; DB 10; Length 319;  
 Best Local Similarity 94.1%; Pred. No. 3.6e+03;  
 Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0:

Qy 1 GGGAGCCCCAGCAATGC 17  
 |||||  
 Db 178 GGGAGCCCCAGCAACGC 162

RESULT 10  
 A1449767/c  
 LOCUS  
 DEFINITION  
 mr69a08.x1 Stratagene mouse testis (#937308) Mus musculus cDNA  
 clone IMAGE:602678 3', mRNA sequence.

ACCESSION  
 A1449767  
 VERSION  
 A1449767.1 GI:4292306  
 KEYWORDS  
 EST.  
 SOURCE  
 house mouse.  
 ORGANISM  
 Mus musculus  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Mus.

REFERENCE  
AUTHORS

1 (bases 1 to 320)  
 Marra,M., Hillier,L., Kucaba,T., Martin,J., Beck,C., Wylie,T.,  
 Underwood,K., Steptoe,M., Theising,B., Allen,M., Bowers,Y., Person  
 B., Swaller,T., Gibbons,M., Pape,D., Harvey,N., Schurk,R., Ritter  
 E., Kohn,S., Shin,T., Jackson,I., Cardenas,M., McCann,R.,  
 Waterston,R. and Wilson,R.  
 The WashU-NCI Mouse EST Project 1999  
 Unpublished (1999)  
 Contact: Marra M/WashU-NCI Mouse EST Project 1999  
 Washington University School of Medicine  
 4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108, USA  
 Tel: 314 286 1800  
 Fax: 314 286 1810  
 Email: mouseest@wustl.edu

TITLE  
JOURNAL  
COMMENT

This clone is available royalty-free through LNL; contact the  
 IMAGE Consortium (info@image.lnl.gov) for further information.  
 This clone was previously sequenced on the 5' end only, this new  
 data is from the 3' end  
 Possible reversed clone: polyT not found.

FEATURES  
source

1..320  
 Location/Qualifiers  
 /organism="Mus musculus"  
 /strain="CD-1"  
 /db\_xref="taxon:10090"  
 /clone="IMAGE:602678"  
 /clone\_lib="Stratagene mouse testis (#937308)"  
 /sex="males"  
 /tissue\_type="testis"  
 /dev\_stage="10-12 week old"  
 /lab\_host="SOLR (kanamycin resistant)"  
 /note="Organ: testis; Vector: pBluescript SK-; Site\_1:  
 EcoRI; Site\_2: XhoI; Cloned unidirectionally. Primer:  
 Oligo dT. Average insert size: 1.0 kb; Uni-ZAP XR Vector;  
 -5' adaptor sequence: 5' GAATTCGGCAGCAG 3' -3' adaptor  
 sequence: 5' CTCGAGTTTTTTTTTTT 3'"

BASE COUNT 81 a 70 c 85 g 84 t  
 ORIGIN  
 Query Match 90.6%; Score 15.4; DB 9; Length 320;  
 Best Local Similarity 94.1%; Pred. No. 3.6e+03;  
 Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0:

Qy 1 GGGAGCCCCAGCAATGC 17  
 |||||  
 Db 34 GGGAGCCCCAGCAATGC 18

RESULT 11  
 AA629118/c  
 LOCUS  
 DEFINITION  
 af57e02.s1 Soares total fetus.Nb2HF8.9w Homo sapiens cDNA clone  
 IMAGE:1035770 3' similar to gb:M23102 HIGH AFFINITY NERVE GROWTH  
 FACTOR RECEPTOR PRECURSOR (HUMAN);, mRNA sequence.

ACCESSION  
 AA629118  
 VERSION  
 AA629118.1 GI:2541505  
 KEYWORDS  
 EST.  
 SOURCE  
 human.  
 ORGANISM  
 Homo sapiens  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE  
 AUTHORS  
 1 (bases 1 to 336)  
 Hillier,L., Allen,M., Bowles,L., Dubuque,T., Geisel,G., Jost,S.,  
 Krizman,D., Kucaba,T., Lacy,M., Le,N., Lennon,G., Marra,M., Martin  
 J., Moore,B., Schellenberg,K., Steptoe,M., Tan,F., Theising,B.,  
 White,Y., Wylie,T., Waterston,R. and Wilson,R.

WashU-NCI human EST Project  
 Unpublished (1997)  
 Contact: Willson RK  
 Washington University School of Medicine  
 4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108  
 Tel: 314 286 1800  
 Fax: 314 286 1810  
 Email: est@wustl.edu

This clone is available royalty-free through LBNL; contact the IMAGE Consortium (info@imga.lbnl.gov) for further information. Trace considered overall poor quality. Insert length: 710 Std Error: 0.00 Seq primer: -40ml3 fwd. FT from Amersham high quality sequence stop: 1. Location/Qualifiers

FEATURES

source

1. 336  
/organism="Homo sapiens"  
/db\_xref="Laxon:9606"  
/clone="IMAGE:1035770"  
/clone\_lib="Soares\_total\_telus\_Nb2Hf8\_9w"  
/dev\_stage="8-9 weeks"  
/lab\_host="DH10B"

/note="Vector: pT773D-Pac (Pharmacia) with a modified polylinker; Site 1: Not 1; Site 2: Eco RI; 1st strand cDNA was prepared from mRNA obtained from pooled 8-9 week (total) fetus material with a Not 1 - oligo(dT) primer [5' TGTTACCAATCTGAAGTGGGAGCGGCGCTTAATTTTCTTTTCTT 3'] Double-stranded cDNA was ligated to Eco RI adaptors (Pharmacia), digested with Not 1 and cloned into the Not 1 and Eco RI sites of the modified pT73 vector. Library went through one round of normalization, and was constructed by Bento Soares and M. Fatima Bonaldo. "

BASE COUNT 64 a 95 c 98 g 79 t  
ORIGIN  
Query Match 90.6%; Score 15.4; DB 9; Length 336;  
Best Local Similarity 94.1%; Pred. No. 3.6e+03;  
Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 GGGAGCCCGACCAATGC 17  
IIIIIIIIIIIIIIII

Db 301 GGGAGCCCGACCAATGC 285

RESULT 12  
BH842232/c  
LOCUS  
DEFINITION 382 bp DNA linear GSS 13-JUN-2002  
TC3-50P20.TP TC3 Trypanosoma cruzi genomic clone TC3-50P20, DNA sequence.

ACCESSION BH842232 GI:21409447  
VERSION  
KEYWORDS  
SOURCE  
ORGANISM Trypanosoma cruzi.  
Trypanosoma cruzi  
Eukaryota; Euklenozoa; Kinetoplastida; Trypanosomatidae;  
Trypanosoma; Schizotrypanum.

REFERENCE 1 (bases 1 to 382)  
Myler,P.J., Aggarwal,G., Fazelinia,G., Mack,J., Marty,A., Munden,H., Nelson,S., Pentony,M., Rinta,J., Robertson,L., Seyler,A., Sisk,E., Stuart,K., Vogt,C., Worthey,E., El-Sayed,N.M., Ghedin,E. and Andersson,B.  
Trypanosoma cruzi CL-Brener TC3 BAC-end sequencing

TITLE  
JOURNAL  
COMMENT  
Unpublished (2001)  
Other\_GSSs: TC3-50P20.TP.1 TC3-50P20.TV  
Contact: Peter Myler  
Seattle Biomedical Research Institute  
4 Nickerson Street, Seattle, WA 98109, USA  
Tel: 206 284 8846  
Fax: 206 284 0313  
Email: myler@sbri.org  
Clones are derived from the Trypanosoma cruzi CL-Brener BAC library TC3. For clone availability, please contact Dr. Bjorn Andersson at Uppsala University (bjorn.andersson@genpat.uu.se).

Seq primer: SP6  
Class: BAC ends.  
Location/Qualifiers

1. 382  
/organism="Trypanosoma cruzi"  
/strain="CL Brener"  
/db\_xref="taxon:5693"  
/clone="TC3-50P20"

/clone\_lib="TC3"  
/note="Vector: pBelOBAC11; Site 1: Hin dIII; Constructed for Uppsala University by Marie-Christine Le Paslier in the Laboratory of Denis Le Paslier at the Centre d'Etude du Polymorphisme Humain (CEPH), Paris, France. Briefly, Trypanosoma cruzi CL-Brener agarose embedded DNA (obtained from Dr. Franco da Silveira) was partially digested with Hin dIII. High molecular weight fragments were ligated in pBelOBAC11 digested with Hin dIII. The average insert size is 100 kb. Total clone coverage: approx. 33 X the haploid genome."

BASE COUNT 88 a 112 c 81 g 101 t  
ORIGIN

Query Match 90.6%; Score 15.4; DB 17; Length 382;  
Best Local Similarity 94.1%; Pred. No. 3.8e+03;  
Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 GGGAGCCCGACCAATGC 17  
IIIIIIIIIIIIIIII

Db 172 GGGAGCCCGACCAATGC 156

RESULT 13  
BH845093/c  
LOCUS  
DEFINITION 382 bp DNA linear GSS 13-JUN-2002  
TC3-50P20.TP.1 TC3 Trypanosoma cruzi genomic clone TC3-50P20, DNA sequence.

ACCESSION BH845093  
VERSION BH845093.1 GI:21415281  
KEYWORDS  
SOURCE  
ORGANISM Trypanosoma cruzi.  
Trypanosoma cruzi  
Eukaryota; Euklenozoa; Kinetoplastida; Trypanosomatidae;  
Trypanosoma; Schizotrypanum.

REFERENCE 1 (bases 1 to 382)  
Myler,P.J., Aggarwal,G., Fazelinia,G., Mack,J., Marty,A., Munden,H., Nelson,S., Pentony,M., Rinta,J., Robertson,L., Seyler,A., Sisk,E., Stuart,K., Vogt,C., Worthey,E., El-Sayed,N.M., Ghedin,E. and Andersson,B.  
Trypanosoma cruzi CL-Brener TC3 BAC-end sequencing

TITLE  
JOURNAL  
COMMENT  
Unpublished (2001)  
Other\_GSSs: TC3-50P20.TP TC3-50P20.TV  
Contact: Peter Myler  
Seattle Biomedical Research Institute  
4 Nickerson Street, Seattle, WA 98109, USA  
Tel: 206 284 8846  
Fax: 206 284 0313  
Email: myler@sbri.org  
Clones are derived from the Trypanosoma cruzi CL-Brener BAC library TC3. For clone availability, please contact Dr. Bjorn Andersson at Uppsala University (bjorn.andersson@genpat.uu.se).

Seq primer: SP6  
Class: BAC ends.  
Location/Qualifiers

1. 382  
/organism="Trypanosoma cruzi"  
/strain="CL Brener"  
/db\_xref="taxon:5693"  
/clone="TC3-50P20"  
/clone\_lib="TC3"  
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BASE COUNT 88 a 112 c 81 g 101 t  
ORIGIN

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Query Match
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Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

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DB 172 GGGAGCCCCCAGCAATGC 156

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LOCUS
DEFINITION CMO-HN0206-090401-813-ell HN0206 Homo sapiens cDNA, mRNA sequence.
ACCESSION BI005357
VERSION BI005357.1 GI:14409431
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 383)
Nagai,M.A., da Silva,W. Jr., Zago,M.A., Bordin,S., Costa,F.F.,
Goldman,G.H., Carvalho,A.F., Matsukuma,A., Baia,G.S., Simpson,D.H.,
Brunstein,A., deOliveira,P.S., Bucher,P., Jongeneel,C.V., O'Hare
M.J., Soares,F., Brentani,R.R., Reis,L.F., de Souza,S.J. and
Simpson,A.J.
Shotgun sequencing of the human transcriptome with ORF expressed
sequence tags
Proc. Natl. Acad. Sci. U.S.A. 97 (7), 3491-3496 (2000)
20202663
Contact: Simpson A.J.G.
Laboratory of Cancer Genetics
Ludwig Institute for Cancer Research
Rua Prof. Antonio Prudente 109, 4 andar, 01509-010, Sao Paulo-SP,
Brazil
Tel: +55-11-2704922
Fax: +55-11-2707001
Email: asimpson@ludwig.org.br
This sequence was derived from the FAPESP/LICK Human Cancer Genome
Project. This entry can be seen in the following URL
(http://www.ludwig.org.br/scripts/gethtml2.pl?tl=CM0-HN0206-
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Seq primer: puc 18 forward
High quality sequence stop: 383.
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derived from ORESTES PCR (U.S. Letters Patent application
No. 196,716 - Ludwig Institute for Cancer Research)
profiles into the pUC 18 vector. Reverse transcription of
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low stringency conditions."
BASE COUNT 74 a 97 c 89 g 123 t
ORIGIN

Query Match
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Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

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RESULT 15
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LOCUS
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sequence.
ACCESSION BI4181
VERSION BI4181.1 GI:2121930
KEYWORDS GSS.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 385)
Adams,M.D., Kelley,J.M., Rounsley,S.R. and Venter,J.C.
Use of a BAC end Sequence Database for Sequence-Ready Map Building
Unpublished (1997)
Other_GSSs: 1054B4.TV
Contact: Mark Adams
Department of Eukaryotic Genomics
The Institute for Genomic Research
9712 Medical Center Dr., Rockville, MD 20850, USA
Tel: 301 838 0200
Fax: 301 838 0208
Email: mdamas@tigr.org
Clones are available from Research Genetics (info@resgen.com). BAC
end search page:
http://www.tigr.org/tdb/hungen/bac_end_search/bac_end_search.html
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Location/Qualifiers
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CalTech Human BAC Library A"
BASE COUNT 80 a 126 c 94 g 85 t
ORIGIN

Query Match
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Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

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Search completed: January 21, 2003, 15:38:03
Job time : 1203.97 secs

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GenCore version 5.1.3  
Copyright (c) 1993 - 2003 Compuen Ltd.

OM nucleic - nucleic search, using sw model

Run on: January 21, 2003, 14:55:32 : Search time 30.6944 Seconds  
(without alignments)  
246.942 Million cell updates/sec

Title: US-09-853-688-35  
Perfect score: 17  
Sequence: 1 gggagccccagcaatgc 17

Scoring table: IDENTITY\_NUC  
Gapop 10.0 , Gapext 1.0

Searched: 393868 seqs, 222934149 residues

Total number of hits satisfying chosen parameters: 787736

Minimum DB seq length: 0  
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Post-processing: Minimum Match 0%  
Maximum Match 100%  
Listing first 45 summaries

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pred. No. is the number of results predicted by chance to have a  
score greater than or equal to the score of the result being printed,  
and is derived by analysis of the total score distribution.

SUMMARIES

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2	17	100.0	3700	10	US-09-853-688-5		Sequence 5, Appli
3	15.4	90.6	468	10	US-09-864-761-5528		Sequence 5528, Ap
4	15.4	90.6	2820	10	US-09-924-859A-4		Sequence 4, Appli
5	15.4	90.6	5086	10	US-09-880-107-3947		Sequence 3947, Ap
6	15.4	90.6	5145	10	US-09-925-299-206		Sequence 206, App
7	15.4	90.6	5416	10	US-09-954-456-786		Sequence 786, App
8	15.4	90.6	5416	10	US-09-880-107-2094		Sequence 2094, Ap
9	15.4	90.6	5432	12	US-10-044-090-22		Sequence 22, Appli
10	15	88.2	1041	9	US-09-738-626-3422		Sequence 3422, Ap
11	15	88.2	1152	9	US-09-738-626-3424		Sequence 3424, Ap
12	15	88.2	1894	10	US-09-126-945B-1		Sequence 1, Appli
13	15	88.2	1884	10	US-09-841-963A-1		Sequence 1, Appli
14	15	88.2	1905	10	US-09-866-356-2		Sequence 2, Appli
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16	14.4	84.7	302	9	US-10-015-219-718		Sequence 718, App
17	14.4	84.7	302	10	US-09-777-564-718		Sequence 718, App
18	14.4	84.7	316	10	US-09-998-598-1147		Sequence 1147, Ap
19	14.4	84.7	698	10	US-09-910-943-311		Sequence 311, App

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22	14.4	84.7	1732	9	US-10-063-547-81	Sequence 81, Appli
23	14.4	84.7	1732	9	US-10-174-590-281	Sequence 281, App
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25	14.4	84.7	1732	12	US-10-006-867-81	Sequence 81, Appli
26	14.4	84.7	1732	12	US-10-052-586-281	Sequence 281, Appli
27	14.4	84.7	1839	9	US-09-798-051-1	Sequence 1, Appli
28	14	82.4	1734	9	US-10-103-511-3	Sequence 3, Appli
29	14	82.4	1734	10	US-09-805-204-3	Sequence 3, Appli
30	14	82.4	4065	10	US-09-808-571A-3	Sequence 1, Appli
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32	14	82.4	9745	10	US-09-764-869-2259	Sequence 2259, Ap
33	14	82.4	12149	10	US-09-764-869-2258	Sequence 2258, Ap
34	14	82.4	29629	12	US-10-135-689-3	Sequence 3, Appli
35	14	82.4	170834	10	US-09-835-232-7	Sequence 7, Appli
36	13.8	81.2	293	10	US-09-983-965-2249	Sequence 2249, Ap
37	13.8	81.2	337	10	US-09-983-965-2463	Sequence 2463, Ap
38	13.8	81.2	349	10	US-09-983-965-2493	Sequence 2493, Ap
39	13.8	81.2	362	10	US-09-983-965-2530	Sequence 2530, Ap
40	13.8	81.2	365	9	US-09-796-692-5576	Sequence 5576, Ap
41	13.8	81.2	379	10	US-09-783-590-7717	Sequence 7717, Ap
42	13.8	81.2	384	9	US-10-040-739-1302	Sequence 1302, Ap
43	13.8	81.2	390	10	US-09-960-352-3031	Sequence 3031, Ap
44	13.8	81.2	398	10	US-09-728-445-656	Sequence 656, App
45	13.8	81.2	479	9	US-09-796-692-3578	Sequence 3578, Ap

ALIGNMENTS

RESULT 1  
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: Sequence 35, Application US/09853688  
: Patent No. US20020081605A1  
: GENERAL INFORMATION:  
: APPLICANT: COOPER, DAVID N.  
: APPLICANT: PROCTER, ANNIE M.  
: APPLICANT: GREGORY, JOHN  
: APPLICANT: MILLAR, DAVID S.  
: TITLE OF INVENTION: METHOD FOR DETECTING GROWTH HORMONE VARIATIONS IN  
: FILE REFERENCE: WCM78  
: CURRENT APPLICATION NUMBER: US/09/853,688  
: CURRENT FILING DATE: 2001-05-14  
: NUMBER OF SEQ ID NOS: 66  
: SOFTWARE: PatentIn Ver. 2.1  
: SEQ ID NO 35  
: LENGTH: 17  
: TYPE: DNA  
: ORGANISM: Homo sapiens  
US-09-853-688-35

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Oy 1 GGGAGCCCCAGCAATGC 17  
Db 1 GGGAGCCCCAGCAATGC 17

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US-09-853-688-5  
: Sequence 5, Application US/09853688  
: Patent No. US20020081605A1  
: GENERAL INFORMATION:  
: APPLICANT: COOPER, DAVID N.  
: APPLICANT: PROCTER, ANNIE M.  
: APPLICANT: GREGORY, JOHN  
: APPLICANT: MILLAR, DAVID S.  
: TITLE OF INVENTION: METHOD FOR DETECTING GROWTH HORMONE VARIATIONS IN  
: FILE REFERENCE: WCM78  
: CURRENT APPLICATION NUMBER: US/09/853,688  
: CURRENT FILING DATE: 2001-05-14  
: NUMBER OF SEQ ID NOS: 66  
: SOFTWARE: PatentIn Ver. 2.1  
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: LENGTH: 17  
: TYPE: DNA  
: ORGANISM: Homo sapiens  
US-09-853-688-5

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; FILE REFERENCE: WCM78
; CURRENT APPLICATION NUMBER: US/09/853,688
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US-09-853-688-5

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; Sequence 5528, Application US/09864761
; Patent No. US20020048763A1
; GENERAL INFORMATION:
; APPLICANT: Penn, Sharon G.
; APPLICANT: Rank, David R.
; APPLICANT: Hanzel, David K.
; APPLICANT: Chen, Wensheng
; TITLE OF INVENTION: HUMAN GENOME-DERIVED SINGLE EXON NUCLEIC ACID PROBES USEFUL FOR
; FILE REFERENCE: Aeomica-X-1
; CURRENT APPLICATION NUMBER: US/09/864,761
; CURRENT FILING DATE: 2001-05-23
; PRIOR APPLICATION NUMBER: US 60/180,312
; PRIOR FILING DATE: 2000-02-04
; PRIOR APPLICATION NUMBER: US 60/207,456
; PRIOR FILING DATE: 2000-05-26
; PRIOR APPLICATION NUMBER: US 09/632,366
; PRIOR FILING DATE: 2000-08-03
; PRIOR APPLICATION NUMBER: GB 24263.6
; PRIOR FILING DATE: 2000-10-04
; PRIOR APPLICATION NUMBER: US 60/236,359
; PRIOR FILING DATE: 2000-09-27
; PRIOR APPLICATION NUMBER: PCT/US01/00666
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00667
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00664
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; PRIOR APPLICATION NUMBER: PCT/US01/00665
; PRIOR FILING DATE: 2001-01-30
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; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: US 60/234,687
; PRIOR FILING DATE: 2000-09-21
; PRIOR APPLICATION NUMBER: US 09/608,408
; PRIOR FILING DATE: 2000-06-30
; PRIOR APPLICATION NUMBER: US 09/774,203
; PRIOR FILING DATE: 2001-01-29
; NUMBER OF SEQ ID NOS: 49117
; SOFTWARE: Annomax Sequence Listing Engine vers. 1.1
; SEQ ID NO 5528

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; OTHER INFORMATION: EXPRESSED IN LUNG, SIGNAL = 1.9
; OTHER INFORMATION: EXPRESSED IN HELA, SIGNAL = 2.1
; OTHER INFORMATION: EXPRESSED IN ADULT LIVER, SIGNAL = 1.7
; OTHER INFORMATION: EXPRESSED IN PLACENTA, SIGNAL = 1.6
; OTHER INFORMATION: EXPRESSED IN HEART, SIGNAL = 1.6
; OTHER INFORMATION: EXPRESSED IN FETAL LIVER, SIGNAL = 2.1
; OTHER INFORMATION: EXPRESSED IN BRAIN, SIGNAL = 1.9
; OTHER INFORMATION: EXPRESSED IN BONE MARROW, SIGNAL = 1.5
US-09-864-761-5528

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; Patent No. US20020137113A1
; GENERAL INFORMATION:
; APPLICANT: Godowski, Paul J.
; APPLICANT: Mark, Melanie R.
; APPLICANT: Sadick, Michael D.
; APPLICANT: Shelton, David L.
; APPLICANT: Wong, Wai Lee Tan
; TITLE OF INVENTION: KINASE RECEPTOR ACTIVATION ASSAY
; FILE REFERENCE: P0854C1P2C1
; CURRENT APPLICATION NUMBER: US/09/924,859A
; CURRENT FILING DATE: 2001-08-08
; PRIOR APPLICATION NUMBER: US/09/417,381
; PRIOR FILING DATE: 1999-10-13
; NUMBER OF SEQ ID NOS: 11
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US-09-924-859A-4

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RESULT 5
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; Sequence 3947, Application US/09880107
; Patent No. US20020142981A1
; GENERAL INFORMATION:
; APPLICANT: Horne, Darci T.
; APPLICANT: Vockley, Joseph G.
; APPLICANT: Scherf, Uwe
; APPLICANT: Gene Logic, Inc.
; TITLE OF INVENTION: Gene Expression Profiles in Liver Cancer
; FILE REFERENCE: 44921-5028-WO
; CURRENT APPLICATION NUMBER: US/09/880,107
; CURRENT FILING DATE: 2001-06-14
; PRIOR APPLICATION NUMBER: US 60/211,379
; PRIOR FILING DATE: 2000-06-14
; PRIOR APPLICATION NUMBER: US 60/237,054
; PRIOR FILING DATE: 2000-10-02

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; TYPE: DNA
; ORGANISM: Homo sapiens
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; OTHER INFORMATION: Genbank Accession No. US20020142981A1 Z74616
; NAME/KEY: unsure
; LOCATION: (1)..(5086)
; OTHER INFORMATION: n = a or c or g or t
US-09-880-107-3947

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Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

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Db 1102 GGGAGCCCCCAGCAACGC 1086

RESULT 6
US-09-925-299-206/c
; Sequence 206, Application US/09925299
; Patent No. US20020055627A1
; GENERAL INFORMATION:
; APPLICANT: Rosen et al.
; TITLE OF INVENTION: Nucleic Acids, Proteins and Antibodies
; FILE REFERENCE: PA102
; CURRENT APPLICATION NUMBER: US/09/925,299
; PRIOR FILING DATE: 2001-08-10
; PRIOR APPLICATION NUMBER: PCT/US00/05883
; PRIOR FILING DATE: 2000-03-08
; PRIOR APPLICATION NUMBER: 60/124,270
; PRIOR FILING DATE: 1999-03-12
; NUMBER OF SEQ ID NOS: 1556
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 206
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; TYPE: DNA
; ORGANISM: Homo sapiens
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; LOCATION: (5126)
; OTHER INFORMATION: n equals a,t,g, or c
; NAME/KEY: misc_feature
; LOCATION: (5143)
; OTHER INFORMATION: n equals a,t,g, or c
US-09-925-299-206

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Best Local Similarity 94.1%; Pred. No. 52;
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Db 1105 GGGAGCCCCCAGCAACGC 1089

RESULT 7
US-09-954-456-786/c
; Sequence 786, Application US/09954456
; Patent No. US20020115057A1
; GENERAL INFORMATION:
; APPLICANT: Young, Paul
; TITLE OF INVENTION: Process for Identifying Anti-Cancer Therapeutic Agents Using Cand
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; TITLE OF INVENTION: Sets
; FILE REFERENCE: 689290-76
; CURRENT APPLICATION NUMBER: US/09/954,456
; PRIOR FILING DATE: 2001-09-18
; PRIOR APPLICATION NUMBER: US/60/233,617
; PRIOR FILING DATE: 2000-09-18
; PRIOR APPLICATION NUMBER: US/60/234,052
; PRIOR FILING DATE: 2000-09-20
; PRIOR APPLICATION NUMBER: US/60/234,923
; PRIOR FILING DATE: 2000-09-25
; PRIOR APPLICATION NUMBER: US/60/235,134
; PRIOR FILING DATE: 2000-09-25
; PRIOR APPLICATION NUMBER: US/60/235,637
; PRIOR FILING DATE: 2000-09-26
; PRIOR APPLICATION NUMBER: US/60/235,638
; PRIOR FILING DATE: 2000-09-26
; PRIOR APPLICATION NUMBER: US/60/235,711
; PRIOR FILING DATE: 2000-09-27
; PRIOR APPLICATION NUMBER: US/60/235,720
; PRIOR FILING DATE: 2000-09-27
; PRIOR APPLICATION NUMBER: US/60/235,840
; PRIOR FILING DATE: 2000-09-27
; PRIOR APPLICATION NUMBER: US/60/235,863
; NUMBER OF SEQ ID NOS: 2276
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 786
; LENGTH: 5416
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-954-456-786

Query Match          90.6%; Score 15.4; DB 10; Length 5416;
Best Local Similarity 94.1%; Pred. No. 52;
Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

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RESULT 8
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; Sequence 2094, Application US/09880107
; Patent No. US20020142981A1
; GENERAL INFORMATION:
; APPLICANT: Horne, Darci T.
; APPLICANT: Vockley, Joseph G.
; APPLICANT: Scherf, Jwe
; APPLICANT: Gene Logic, Inc.
; TITLE OF INVENTION: Gene Expression Profiles in Liver Cancer
; FILE REFERENCE: 44921-5028-WO
; CURRENT APPLICATION NUMBER: US/09/880,107
; CURRENT FILING DATE: 2001-06-14
; PRIOR APPLICATION NUMBER: US 60/211,379
; PRIOR FILING DATE: 2000-06-14
; PRIOR APPLICATION NUMBER: US 60/237,054
; PRIOR FILING DATE: 2000-10-02
; NUMBER OF SEQ ID NOS: 3950
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 2094
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; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; OTHER INFORMATION: Genbank Accession No. US20020142981A1 J03464
US-09-880-107-2094

Query Match          90.6%; Score 15.4; DB 10; Length 5416;
Best Local Similarity 94.1%; Pred. No. 52;
Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 GGGAGCCCCCAGCAATGC 17
```

```

DB 1412 GGGAGCCCCCAGCAATGC 1416
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RESULT 9
US-10-044-090-22/c
; Sequence 22, Application US/10044090
; Patent No. US20020137081A1
; GENERAL INFORMATION:
; APPLICANT: Olga Bandman
; TITLE OF INVENTION: GENES DIFFERENTIALLY EXPRESSED IN VASCULAR TISSUE ACTIVATION
; FILE REFERENCE: PA-0028 US
; CURRENT APPLICATION NUMBER: US/10/044,090
; CURRENT FILING DATE: 2002-01-09
; NUMBER OF SEQ ID NOS: 850
; SOFTWARE: PERL Program
; SEQ ID NO 22
; LENGTH: 5432
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc feature
; OTHER INFORMATION: Incyte ID No. US20020137081A1 1383093.13
US-10-044-090-22
Query Match 90.68; Score 15.4; DB 12: Length 5432;
Best Local Similarity 94.1%; Pred. No. 52: 1; Indels 0; Gaps 0;
Matches 16; Conservative 0; Mismatches 0;

QY 1 GGGAGCCCCCAGCAATGC 17
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DB 1435 GGGAGCCCCCAGCAATGC 1419

RESULT 10
US-09-738-626-3422
; Sequence 3422, Application US/09738626
; Publication No. US20020197605A1
; GENERAL INFORMATION:
; APPLICANT: NAKAGAWA, SATOSHI
; APPLICANT: ANDO, SEIKO
; APPLICANT: HAYASHI, MIKIRO
; APPLICANT: OCHIAI, KEIKO
; APPLICANT: YOKOI, HARUHIKO
; APPLICANT: TATEISHI, NAOKO
; APPLICANT: SENOH, AKIHIRO
; APPLICANT: IKEDA, MASATO
; APPLICANT: OZAKI, AKIO
; TITLE OF INVENTION: NOVEL POLYNUCLEOTIDES
; FILE REFERENCE: 249-125
; CURRENT APPLICATION NUMBER: US/09/738,626
; CURRENT FILING DATE: 2000-12-18
; PRIOR APPLICATION NUMBER: JP 99/377484
; PRIOR FILING DATE: 1999-12-16
; PRIOR APPLICATION NUMBER: JP 00/159162
; PRIOR FILING DATE: 2000-04-07
; PRIOR APPLICATION NUMBER: JP 00/280988
; PRIOR FILING DATE: 2000-08-03
; NUMBER OF SEQ ID NOS: 7059
; SOFTWARE: PatentIn ver. 3.0
; SEQ ID NO 3422
; LENGTH: 1041
; TYPE: DNA
; ORGANISM: Corynebacterium glutamicum
US-09-738-626-3422
Query Match 88.2%; Score 15; DB 9: Length 1041;
Best Local Similarity 100.0%; Pred. No. 74;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 GAGCCCCCAGCAATGC 17
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DB 496 GAGCCCCCAGCAATGC 510
RESULT 11
US-09-738-626-3424/c
; Sequence 3424, Application US/09738626
; Publication No. US20020197605A1
; GENERAL INFORMATION:
; APPLICANT: NAKAGAWA, SATOSHI
; APPLICANT: MIZOGUCHI, HIROSHI
; APPLICANT: ANDO, SEIKO
; APPLICANT: HAYASHI, MIKIRO
; APPLICANT: OCHIAI, KEIKO
; APPLICANT: YOKOI, HARUHIKO
; APPLICANT: TATEISHI, NAOKO
; APPLICANT: SENOH, AKIHIRO
; APPLICANT: IKEDA, MASATO
; APPLICANT: OZAKI, AKIO
; TITLE OF INVENTION: NOVEL POLYNUCLEOTIDES
; FILE REFERENCE: 249-125
; CURRENT APPLICATION NUMBER: US/09/738,626
; CURRENT FILING DATE: 2000-12-18
; PRIOR APPLICATION NUMBER: JP 99/377484
; PRIOR FILING DATE: 1999-12-16
; PRIOR APPLICATION NUMBER: JP 00/159162
; PRIOR FILING DATE: 2000-04-07
; PRIOR APPLICATION NUMBER: JP 00/280988
; PRIOR FILING DATE: 2000-08-03
; NUMBER OF SEQ ID NOS: 7059
; SOFTWARE: PatentIn ver. 3.0
; SEQ ID NO 3424
; LENGTH: 1152
; TYPE: DNA
; ORGANISM: Corynebacterium glutamicum
US-09-738-626-3424
Query Match 88.2%; Score 15; DB 9: Length 1152;
Best Local Similarity 100.0%; Pred. No. 74;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 GAGCCCCCAGCAATGC 17
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DB 858 GAGCCCCCAGCAATGC 844

RESULT 12
US-09-126-945B-1
; Sequence 1, Application US/09126945B
; Patent No. US20010010934A1
; GENERAL INFORMATION:
; APPLICANT: Libermann, Towia A.
; APPLICANT: Oeltgen, Joerg P.
; APPLICANT: Kunsch, Charles A.
; APPLICANT: Endress, Gregory A.
; APPLICANT: Rosen, Craig A.
; TITLE OF INVENTION: Prostate Derived Bts Factor
; FILE REFERENCE: 1486.1090000
; CURRENT APPLICATION NUMBER: US/09/126,945B
; CURRENT FILING DATE: 1998-07-31
; NUMBER OF SEQ ID NOS: 15
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 1
; LENGTH: 1894
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-126-945B-1
Query Match 88.2%; Score 15; DB 10: Length 1894;
Best Local Similarity 100.0%; Pred. No. 77;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 GGAGCCCCCAGCAATGC 16
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100 864 GGAGCCCCAGCAATG 878

RESULT 13
US-09-841-963A-1
: Sequence 1, Application US/09841963A
: Patent No. US20020081601A1
: GENERAL INFORMATION:
: APPLICANT: Watson, Dennis K.
: APPLICANT: Papas, Takis S. (Deceased)
: APPLICANT: Papas, Tula C. (Legal Representative)
: TITLE OF INVENTION: Methods and compositions for the diagnosis and treatment of cancer
: TITLE OF INVENTION: based on transcription factor ETS2
: FILE REFERENCE: 10545-015-999
: CURRENT APPLICATION NUMBER: US/09/841,963A
: CURRENT FILING DATE: 2001-04-25
: PRIOR APPLICATION NUMBER: PCT/US99/27805
: PRIOR FILING DATE: 1999-11-23
: PRIOR APPLICATION NUMBER: 06/109,850
: PRIOR FILING DATE: 1998-11-25
: NUMBER OF SEQ ID NOS: 6
: SOFTWARE: PatentIn version 3.0
: SEQ ID NO 1
: LENGTH: 1894
: TYPE: DNA
: ORGANISM: Homo sapiens
: FEATURE:
: NAME/KEY: CDS
: LOCATION: (416)..(1423)
US-09-841-963A-1

Query Match
Best Local Similarity 100.0%; Pred. No. 77; Length 1894;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 GGAGCCCCAGCAATG 16
| | | | | | | | | | | | | | | | | | | | | |
Db 864 GGAGCCCCAGCAATG 878

RESULT 14
US-09-866-356-2
: Sequence 2, Application US/09866356
: Patent No. US20020098543A1
: GENERAL INFORMATION:
: APPLICANT: Bandman, Olga
: Corley, Neil C.
: Guegler, Karl J.
: Lal, Preeti
: TITLE OF INVENTION: PROSTATE-ASSOCIATED ETS PROTEIN
: NUMBER OF SEQUENCES: 4
: CORRESPONDENCE ADDRESS:
: ADDRESSEE: Incyte Pharmaceuticals, Inc.
: STREET: 3174 Porter Dr.
: CITY: Palo Alto
: STATE: CA
: COUNTRY: USA
: ZIP: 94304
: COMPUTER READABLE FORM:
: MEDIUM TYPE: Diskette
: COMPUTER: IBM Compatible
: OPERATING SYSTEM: DOS
: SOFTWARE: FastSeq for Windows Version 2.0
: CURRENT APPLICATION DATA:
: APPLICATION NUMBER: US/09/866,356
: FILING DATE: 29-May-2001
: PRIOR APPLICATION DATA:
: APPLICATION NUMBER: 09/055,113
: FILING DATE: 1998-04-03
: ATTORNEY/AGENT INFORMATION:
: NAME: Billings, Lucy J.
: REGISTRATION NUMBER: 36,749
: REFERENCE/DOCKET NUMBER: PP-0501 US
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: TELECOMMUNICATION INFORMATION:
: TELEPHONE: 650-855-0555
: TELEFAX: 650-845-4166
: TELEX: <Unknown>
: INFORMATION FOR SEQ ID NO: 2:
: SEQUENCE CHARACTERISTICS:
: LENGTH: 1905 base pairs
: TYPE: nucleic acid
: STRANDEDNESS: single
: TOPOLOGY: linear
: IMMEDIATE SOURCE:
: LIBRARY: PROSTUT12
: CLONE: 1813005
: SEQUENCE DESCRIPTION: SEQ ID NO: 2:
US-09-866-356-2

Query Match 88.2%; Score 15; DB 10; Length 1905;
Best Local Similarity 100.0%; Pred. No. 77;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 GGAGCCCCAGCAATG 16
| | | | | | | | | | | | | | | | | | | | | |
Db 878 GGAGCCCCAGCAATG 892

RESULT 15
US-09-962-436-91
: Sequence 91, Application US/09962436
: Patent No. US20020081301A1
: GENERAL INFORMATION:
: APPLICANT: Soppet, Daniel
: TITLE OF INVENTION: Cancer Gene Determination and Therapeutic Screening Using S
: FILE REFERENCE: 689290-75
: CURRENT APPLICATION NUMBER: US/09/962,436
: CURRENT FILING DATE: 2001-09-25
: PRIOR APPLICATION NUMBER: US/60/235,082
: PRIOR FILING DATE: 2000-09-25
: PRIOR APPLICATION NUMBER: US/60/234,924
: PRIOR FILING DATE: 2000-09-25
: NUMBER OF SEQ ID NOS: 568
: SOFTWARE: PatentIn version 3.0
: SEQ ID NO 91
: LENGTH: 266
: TYPE: DNA
: ORGANISM: Homo sapiens
: FEATURE:
: NAME/KEY: misc_feature
: OTHER INFORMATION: n=a,t,g or c
US-09-962-436-91

Query Match 84.7%; Score 14.4; DB 10; Length 266;
Best Local Similarity 93.8%; Pred. No. 1,3e+02;
Matches 15; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 2 GGAGCCCCAGCAATGC 17
| | | | | | | | | | | | | | | | | | | | | |
Db 170 GGAGCTCCAGCAATGC 185

Search completed: January 21, 2003, 16:29:15
Job time : 30.6944 secs
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GenCore version 5.1.3  
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CM nucleic - nucleic search, using sw model

Run on: January 21, 2003, 15:09:37 ; Search time 29.2778 seconds  
(without alignments)  
178.070 Million cell updates/sec

Title: US-09-853-688-35

Perfect score: 17

Sequence: 1 gggagcccgcaatgc 17

Scoring table: IDENTITY\_NUC

Gapop 10.0 , Gapext 1.0

Searched: 441362 seqs, 153338381 residues

Total number of hits satisfying chosen parameters: 882724

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Listing first 45 summaries

Database : Issued\_Patents\_NA:\*

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3: /cgn2\_6/ptodata/2/ina/6A\_COMB.seq:\*

4: /cgn2\_6/ptodata/2/ina/6B\_COMB.seq:\*

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6: /cgn2\_6/ptodata/2/ina/backfilesl.seq:\*

Prod. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match %	Length	DB ID	Description
1	15.4	90.6	2820	1	US-08-286-305A-4 Sequence 4, Appli
2	15.4	90.6	2820	2	US-08-441-104A-4 Sequence 4, Appli
3	15.4	90.6	2820	2	US-08-440-816A-4 Sequence 4, Appli
4	15.4	90.6	2820	4	US-09-417-381A-4 Sequence 4, Appli
5	15.4	90.6	38682	4	US-08-943-731-2 Sequence 2, Appli
6	15	88.2	1905	4	US-09-055-113-2 Sequence 2, Appli
7	15	88.2	1929	2	US-09-016-000-10 Sequence 10, Appli
8	14.4	84.7	23	2	US-08-468-551-9 Sequence 9, Appli
9	14.4	84.7	104	3	US-08-717-294-104 Sequence 104, App
10	14.4	84.7	4451	3	US-08-717-294-42 Sequence 42, Appli
11	14.4	84.7	9179	4	US-09-453-702B-162 Sequence 162, App
12	14.4	84.7	12537	2	US-08-611-280-4 Sequence 4, Appli
13	14.4	84.7	12537	4	US-09-195-940-4 Sequence 4, Appli
14	14.4	84.7	12537	4	US-09-562-466-4 Sequence 4, Appli
15	14	82.4	1209	4	US-08-822-774-18 Sequence 18, Appli
16	14	82.4	1209	4	US-09-632-711-18 Sequence 18, Appli
17	14	82.4	1209	4	US-09-632-703B-18 Sequence 18, Appli
18	14	82.4	1209	4	US-09-632-702-18 Sequence 18, Appli
19	14	82.4	7379	4	US-09-341-587-5 Sequence 5, Appli
20	14	82.4	29629	4	US-09-729-995-3 Sequence 3, Appli
21	13.8	81.2	766	4	US-09-105-839P-1 Sequence 1, Appli
22	13.8	81.2	2062	1	US-08-050-319B-24 Sequence 24, Appli
23	13.8	81.2	2062	2	US-08-465-982-24 Sequence 24, Appli
24	13.8	81.2	2081	2	US-09-096-982-7 Sequence 7, Appli
25	13.8	81.2	2081	2	US-08-653-650A-7 Sequence 1, Appli
26	13.8	81.2	2161	3	US-09-106-038A-1 Sequence 3, Appli
27	13.8	81.2	2161	4	US-09-505-250-3 Sequence 3, Appli

28	13.8	81.2	2175	1	US-08-321-668-1 Sequence 1, Appli
29	13.8	81.2	2175	1	US-08-837-941-1 Sequence 1, Appli
30	13.8	81.2	2175	1	US-08-126-016-1 Sequence 1, Appli
31	13.8	81.2	2175	4	US-08-084-970-1 Sequence 1, Appli
32	13.8	81.2	2230	4	US-09-189-527-12 Sequence 12, Appli
33	13.8	81.2	2834	4	US-09-305-384-6 Sequence 6, Appli
34	13.8	81.2	3196	2	US-09-096-982-4 Sequence 4, Appli
35	13.8	81.2	3196	2	US-08-653-650A-4 Sequence 4, Appli
36	13.8	81.2	6235	4	US-09-305-384-5 Sequence 5, Appli
37	13.8	81.2	6679	4	US-09-305-384-1 Sequence 1, Appli
38	13.4	78.8	189	1	US-08-473-981A-1 Sequence 1, Appli
39	13.4	78.8	189	2	US-08-474-087-1 Sequence 1, Appli
40	13.4	78.8	680	4	US-08-943-731-92 Sequence 92, Appli
41	13.4	78.8	777	4	US-08-998-416-247 Sequence 247, App
42	13.4	78.8	1491	4	US-09-662-249A-3 Sequence 3, Appli
43	13.4	78.8	1781	1	US-08-314-615-2 Sequence 2, Appli
44	13.4	78.8	1781	1	US-08-314-362-2 Sequence 2, Appli
45	13.4	78.8	1781	1	US-08-433-010-2 Sequence 2, Appli

ALIGNMENTS

RESULT 1  
US-08-286-305A-4  
; Sequence 4, Application US/08286305A  
; Patent No. 5766863  
; GENERAL INFORMATION:  
; APPLICANT: Godowski, Paul J.  
; APPLICANT: Mark, Melanie R.  
; APPLICANT: Sadick, Michael D.  
; APPLICANT: Shelton, David L.  
; APPLICANT: Wong, Wai Lee Tan  
; TITLE OF INVENTION: KINASE RECEPTOR ACTIVATION ASSAY  
; NUMBER OF SEQUENCES: 11  
; CORRESPONDENCE ADDRESS:  
; ADDRESSEE: Genentech, Inc.  
; STREET: 460 Point San Bruno Blvd  
; CITY: South San Francisco  
; STATE: California  
; COUNTRY: USA  
; ZIP: 94080

COMPUTER READABLE FORM:  
; MEDIUM TYPE: 5.25 inch, 360 Kb floppy disk  
; COMPUTER: IBM PC compatible  
; OPERATING SYSTEM: PC-DOS/MS-DOS  
; SOFTWARE: patin (Genentech)  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/08/286.305A  
; FILING DATE: 05-AUG-1994  
; CLASSIFICATION: 435  
; PRIOR APPLICATION DATA:  
; APPLICATION NUMBER: 08/170558  
; FILING DATE: 20-DEC-1993  
; PRIOR APPLICATION DATA:  
; APPLICATION NUMBER: 08/157563  
; FILING DATE: 23-NOV-1993  
; ATTORNEY/AGENT INFORMATION:  
; NAME: Lee, Wendy M.  
; REGISTRATION NUMBER: 00.000  
; REFERENCE/DOCKET NUMBER: 854C1P1  
; TELECOMMUNICATION INFORMATION:  
; TELEPHONE: 415/225-1994  
; TELEFAX: 415/952-9881  
; TELEX: 910/371-7168  
; INFORMATION FOR SEQ ID NO: 4:  
; SEQUENCE CHARACTERISTICS:  
; LENGTH: 2820 bases  
; TYPE: nucleic acid  
; STRANDEDNESS: double  
; TOPOLOGY: linear

US-08-286-305A-4

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Query Match          90.6%; Score 15.4; DB 1; Length 2820;
Best Local Similarity 94.1%; Pred. No. 40;
Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 GGGAGCCCCAGCAATGC 17
    |||||
Db 2643 GGGAGCCCCAGCAACGC 2659

RESULT 2
US-08-441-104A-4
; Sequence 4, Application US/08441104A
; Patent No. 5891650
; GENERAL INFORMATION:
; APPLICANT: Godowski, Paul J.
; APPLICANT: Mark, Melanie R.
; APPLICANT: Sadick, Michael D.
; APPLICANT: Shelton, David L.
; APPLICANT: Wong, Wai Lee Tan
; TITLE OF INVENTION: KINASE RECEPTOR ACTIVATION ASSAY
; NUMBER OF SEQUENCES: 11
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Genentech, Inc.
; STREET: 460 Point San Bruno Blvd
; CITY: South San Francisco
; STATE: California
; COUNTRY: USA
; ZIP: 94080
; COMPUTER READABLE FORM:
; MEDIUM TYPE: 3.5 inch, 720 kb floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: patin (Genentech)
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/441,104A
; FILING DATE: 15-MAY-1995
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/286305
; FILING DATE: 05-AUG-1994
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/170558
; FILING DATE: 20-DEC-1993
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/157563
; FILING DATE: 23-NOV-1993
; ATTORNEY/AGENT INFORMATION:
; NAME: Lee, Wendy M.
; REGISTRATION NUMBER: 00,000
; REFERENCE/DOCKET NUMBER: 854C1PIC2
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 415/225-1994
; TELEFAX: 415/952-9881
; TELEX: 910/371-7168
; INFORMATION FOR SEQ ID NO: 4:
; LENGTH: 2820 bases
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
US-08-441-104A-4

Query Match          90.6%; Score 15.4; DB 2; Length 2820;
Best Local Similarity 94.1%; Pred. No. 40;
Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 GGGAGCCCCAGCAATGC 17
    |||||
Db 2643 GGGAGCCCCAGCAACGC 2659

RESULT 3
US-08-440-816A-4
; Sequence 4, Application US/09417381A
; Patent No. 6287784
; GENERAL INFORMATION:
; APPLICANT: Godowski, Paul J.
; APPLICANT: Mark, Melanie R.
; APPLICANT: Sadick, Michael D.
; APPLICANT: Shelton, David L.
; APPLICANT: Wong, Wai Lee Tan
; TITLE OF INVENTION: KINASE RECEPTOR ACTIVATION ASSAY
; FILE REFERENCE: P0854C1P2C1
; CURRENT APPLICATION NUMBER: US/09/417,381A
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; Sequence 4, Application US/08440816A
; Patent No. 5914237
; GENERAL INFORMATION:
; APPLICANT: Godowski, Paul J.
; APPLICANT: Mark, Melanie R.
; APPLICANT: Sadick, Michael D.
; APPLICANT: Shelton, David L.
; APPLICANT: Wong, Wai Lee Tan
; TITLE OF INVENTION: KINASE RECEPTOR ACTIVATION ASSAY
; NUMBER OF SEQUENCES: 11
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Genentech, Inc.
; STREET: 460 Point San Bruno Blvd
; CITY: South San Francisco
; STATE: California
; COUNTRY: USA
; ZIP: 94080
; COMPUTER READABLE FORM:
; MEDIUM TYPE: 5.25 inch, 360 kb floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: patin (Genentech)
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/440,816A
; FILING DATE: 15-MAY-1995
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/286305
; FILING DATE: 05-AUG-1994
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/170558
; FILING DATE: 20-DEC-1993
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/157563
; FILING DATE: 23-NOV-1993
; ATTORNEY/AGENT INFORMATION:
; NAME: Lee, Wendy M.
; REGISTRATION NUMBER: 00,000
; REFERENCE/DOCKET NUMBER: 854C1PIC3
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 415/225-1994
; TELEFAX: 415/952-9881
; TELEX: 910/371-7168
; INFORMATION FOR SEQ ID NO: 4:
; LENGTH: 2820 bases
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
US-08-440-816A-4

Query Match          90.6%; Score 15.4; DB 2; Length 2820;
Best Local Similarity 94.1%; Pred. No. 40;
Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 GGGAGCCCCAGCAATGC 17
    |||||
Db 2643 GGGAGCCCCAGCAACGC 2659

RESULT 4
US-09-417-381A-4
; Sequence 4, Application US/09417381A
; Patent No. 6287784
; GENERAL INFORMATION:
; APPLICANT: Godowski, Paul J.
; APPLICANT: Mark, Melanie R.
; APPLICANT: Sadick, Michael D.
; APPLICANT: Shelton, David L.
; APPLICANT: Wong, Wai Lee Tan
; TITLE OF INVENTION: KINASE RECEPTOR ACTIVATION ASSAY
; FILE REFERENCE: P0854C1P2C1
; CURRENT APPLICATION NUMBER: US/09/417,381A
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; CURRENT FILING DATE: 1999-10-13  
; NUMBER OF SEQ ID NOS: 11  
; SEQ ID NO 4  
; LENGTH: 2820  
; TYPE: DNA  
; ORGANISM: Homo Sapien  
US-09-417-381A-4

Query Match 90.6%; Score 15.4; DB 4; Length 2820;  
Best local Similarity 94.1%; Pred. No. 40;  
Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 GGGAGCCCCAGCAATGC 17  
|||||

DB 2643 GGGAGCCCCAGCAACGC 2659

## RESULT 5

US-08-943-731-2/c  
; Sequence 2, Application US/08943731

; Patent No. 6265157

; GENERAL INFORMATION:

; APPLICANT: PROCKOP, DARWIN J.

; APPLICANT: SPOTILA, LORETTA D.

; APPLICANT: DELTAS, CONSTANTINOS D.

; APPLICANT: SEREDA, LARISSA W.

; APPLICANT: LARSON, MICHAEL

; APPLICANT: PACK, MICHAEL

; APPLICANT: COLIGE, ALAIN

; APPLICANT: EARLY, JAMES

; APPLICANT: KORRKO, JARMO

; APPLICANT: ALA-KORRKO, LEENA, et al.

; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR DETECTING

; TITLE OF INVENTION: ALTERED TYPE I OR TYPE IX COLLAGEN GENE SEQUENCES

; NUMBER OF SEQUENCES: 666

; CORRESPONDENCE ADDRESS:

; ADDRESSEE: PANITCH SCHWARZE JACOBS & NADEL, P.C.

; STREET: ONE COMMERCE SQUARE, 2005 MARKET STREET, 22ND

; CITY: FLR

; CITY: PHILADELPHIA

; STATE: PA

; COUNTRY: USA

; ZIP: 19103-7086

; COMPUTER READABLE FORM:

; MEDIUM TYPE: Floppy disk

; COMPUTER: IBM PC compatible

; OPERATING SYSTEM: PC-DOS/MS-DOS

; SOFTWARE: PatentIn Release #1.0, Version #1.30

; CURRENT APPLICATION DATA:

; APPLICATION NUMBER: US/08/943,731

; FILING DATE: 03-OCT-1997

; CLASSIFICATION: 435

; PRIOR APPLICATION DATA:

; APPLICATION NUMBER: US 08/212,322

; FILING DATE: 14-MAR-1994

; PRIOR APPLICATION DATA:

; APPLICATION NUMBER: US 07/803,628

; FILING DATE: 03-DEC-1991

; ATTORNEY/AGENT INFORMATION:

; NAME: DOYLE LEARY Ph.D., KATHRYN

; REGISTRATION NUMBER: 36,317

; REFERENCE/DOCKET NUMBER: 9598-27

; TELECOMMUNICATION INFORMATION:

; TELEPHONE: 215-965-1284

; TELEFAX: 215-567-2991

; TELEX: 831-494

; INFORMATION FOR SEQ ID NO: 2:

; SEQUENCE CHARACTERISTICS:

; LENGTH: 38682 base pairs

; TYPE: nucleic acid

; STRANDEDNESS: double

; TOPOLOGY: linear

; MOLECULE TYPE: DNA (genomic)

US-08-943-731-2

Query Match 90.6%; Score 15.4; DB 4; Length 38682;  
Best Local Similarity 94.1%; Pred. No. 48;  
Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 GGGAGCCCCAGCAATGC 17  
|||||

DB 17119 GGGAGCCCCAGCAACGC 17103

## RESULT 6

US-09-055-113-2

; Sequence 2, Application US/09055113

; Patent No. 6265565

; GENERAL INFORMATION:

; APPLICANT: Bandman, Olga

; APPLICANT: Corley, Neil C.

; APPLICANT: Guegler, Karl J.

; APPLICANT: Lal, Preeti

; TITLE OF INVENTION: PROSTATE-ASSOCIATED ETS PROTEIN

; NUMBER OF SEQUENCES: 4

; CORRESPONDENCE ADDRESS:

; ADDRESSEE: Incyte Pharmaceuticals, Inc.

; STREET: 3174 Porter Dr.

; CITY: Palo Alto

; STATE: CA

; COUNTRY: USA

; ZIP: 94304

; COMPUTER READABLE FORM:

; MEDIUM TYPE: Diskette

; COMPUTER: IBM Compatible

; OPERATING SYSTEM: DOS

; SOFTWARE: FastSeq for Windows Version 2.0

; CURRENT APPLICATION DATA:

; APPLICATION NUMBER: US/09/055,113

; FILING DATE: Filed Herewith

; PRIOR APPLICATION DATA:

; APPLICATION NUMBER:

; FILING DATE:

; ATTORNEY/AGENT INFORMATION:

; NAME: Billings, Lucy J.

; REGISTRATION NUMBER: 36,749

; REFERENCE/DOCKET NUMBER: PF-0501 US

; TELECOMMUNICATION INFORMATION:

; TELEPHONE: 650-855-0555

; TELEFAX: 650-845-4166

; TELEX:

; INFORMATION FOR SEQ ID NO: 2:

; SEQUENCE CHARACTERISTICS:

; LENGTH: 1905 base pairs

; TYPE: nucleic acid

; STRANDEDNESS: single

; TOPOLOGY: linear

; IMMEDIATE SOURCE:

; LIBRARY: PROSTUT12

; CLONE: 1813005

; US-09-055-113-2

Query Match 88.2%; Score 15; DB 4; Length 1905;

Best Local Similarity 100.0%; Pred. No. 62;

Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 GGAGCCCCAGCAATG 16

|||||

DB 878 GGAGCCCCAGCAATG 892

## RESULT 7

US-09-016-000-10/c

; Sequence 10, Application US/09016000

; Patent No. 5962232

; GENERAL INFORMATION:



APPLICANT: Hillman, Jennifer L.  
APPLICANT: Lal, Preeti  
APPLICANT: Bandman, Olga  
APPLICANT: Akerblom, Ingrid E.  
APPLICANT: Shah, Purvi  
APPLICANT: Corley, Neil C.  
APPLICANT: Guegler, Karl G.  
TITLE OF INVENTION: PROTEIN KINASE MOLECULES  
NUMBER OF SEQUENCES: 12  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Incyte Pharmaceuticals, Inc.  
STREET: 3174 Porter Drive  
CITY: Palo Alto  
STATE: CA  
COUNTRY: USA  
ZIP: 94304

COMPUTER READABLE FORM:  
MEDIUM TYPE: Diskette  
COMPUTER: IBM Compatible  
OPERATING SYSTEM: DOS  
SOFTWARE: FastSeq for Windows Version 2.0  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/09/016,000  
FILING DATE: HEREWITH

CLASSIFICATION:  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER:  
FILING DATE:

ATTORNEY/AGENT INFORMATION:  
NAME: Billings, Lucy J.  
REGISTRATION NUMBER: 36,749  
REFERENCE/DOCKET NUMBER: PF-0465 US  
TELEPHONE: 650-855-0555  
TELEFAX: 650-845-4166  
TELEX:

INFORMATION FOR SEQ ID NO: 10:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 1929 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: single  
TOPOLOGY: linear  
IMMEDIATE SOURCE:  
LIBRARY: MMLRDT01  
CLONE: 472480  
US-09-016-000-10

Query Match 88.2%; Score 15; nb 2; Length 1929;  
Best Local Similarity 100.0%; Pred. No. 62;  
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 GGAGCCCCAGCAATG 16  
|||||

Db 40 GGAGCCCCAGCAATG 26

RESULT 8  
US-08-468-551-9  
Sequence 9, Application US/08468551  
Patent No. 5874212  
GENERAL INFORMATION:  
APPLICANT: Prokop, Darwin J.  
APPLICANT: Rock, Matthew J.  
TITLE OF INVENTION: DETECTION OF SINGLE BASE MUTATIONS AND  
OTHER VARIATIONS IN DOUBLE STRANDED DNA BY  
TITLE OF INVENTION: CONFORMATION-SENSITIVE CELL ELECTROPHORESIS  
NUMBER OF SEQUENCES: 9  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: PANITCH SCHWARZE JACOBS & NADEL, P.C.  
STREET: ONE COMMERCE SQUARE, 2005 MARKET STREET, 22ND  
STREET: FLOOR  
CITY: PHILADELPHIA

STATE: PENNSYLVANIA  
COUNTRY: UNITED STATES  
ZIP: 19103-7086  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC Compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: PatentIn Release #1.0, Version #1.30  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/468,551  
FILING DATE: 06-JUN-1995  
CLASSIFICATION: 435  
ATTORNEY/AGENT INFORMATION:  
NAME: Doyle Leary Ph.D., Kathryn  
REGISTRATION NUMBER: 36,317  
REFERENCE/DOCKET NUMBER: 9855-501  
TELEPHONE: 215-965-1284  
TELEFAX: 215-567-2991  
TELEX: 831-494  
INFORMATION FOR SEQ ID NO: 9:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 23 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: single  
TOPOLOGY: linear  
MOLECULE TYPE: DNA (genomic)  
US-08-468-551-9

Query Match 84.7%; Score 14.4; DB 2; Length 23;  
Best Local Similarity 93.8%; Pred. No. 92;  
Matches 15; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 2 GGAGCCCCAGCAATGC 17  
|||||

Db 8 GGAGCCCCAGCAATGC 23

RESULT 9  
US-08-717-294-104/C  
Sequence 104, Application US/08717294  
Patent No. 6114148  
GENERAL INFORMATION:  
APPLICANT: SEED, BRIAN  
APPLICANT: HAAS, JURGEN  
TITLE OF INVENTION: HIGH LEVEL EXPRESSION OF  
TITLE OF INVENTION: PROTEINS  
NUMBER OF SEQUENCES: 110  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Clark & Elbing LLP  
STREET: 176 Federal Street  
CITY: Boston  
STATE: MA  
COUNTRY: USA  
ZIP: 02110  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Diskette  
COMPUTER: IBM Compatible  
OPERATING SYSTEM: DOS  
SOFTWARE: FastSeq for Windows Version 2.0  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/717,294  
FILING DATE: 20-SEP-1996  
CLASSIFICATION: 435  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER:  
FILING DATE:  
ATTORNEY/AGENT INFORMATION:  
NAME: Elbing, Karen L.  
REGISTRATION NUMBER: 35,238  
REFERENCE/DOCKET NUMBER: 00786/345001  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: 617-428-0200

TELEFAX: 617-428-7045  
TELEX:  
INFORMATION FOR SEQ ID NO: 104:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 104 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: single  
TOPOLOGY: linear  
MOLECULE TYPE: Other  
US-08-717-294-104

Query Match 84.7%; Score 14.4; DB 3; Length 104;  
Best Local Similarity 93.8%; Pred. No. 1.3e+02;  
Matches 15; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 2 GGAGCCCCAGCAATGC 17  
|||||  
Db 98 GGAGCCCCAGCAAGC 83

RESULT 10  
US-08-717-294-42  
Sequence 42, Application US/08717294  
Patent No. 6114148  
GENERAL INFORMATION:  
APPLICANT: SEED, BRIAN  
APPLICANT: HAAS, JURGEN  
TITLE OF INVENTION: HIGH LEVEL EXPRESSION OF  
TITLE OF INVENTION: PROTEINS  
NUMBER OF SEQUENCES: 110  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Clark & Elbing LLP  
STREET: 176 Federal Street  
CITY: Boston  
STATE: MA  
COUNTRY: USA  
ZIP: 02110  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Diskette  
COMPUTER: IBM Compatible  
OPERATING SYSTEM: DOS  
SOFTWARE: FASTSEQ for Windows version 2.0  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/717,294  
FILING DATE: 20-SEP-1996  
CLASSIFICATION: 435  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER:  
FILING DATE:  
ATTORNEY/AGENT INFORMATION:  
NAME: Elbing, Karen L.  
REGISTRATION NUMBER: 35,238  
REFERENCE/DOCKET NUMBER: 00786/345001  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: 617-428-0200  
TELEFAX: 617-428-7045  
TELEX:  
INFORMATION FOR SEQ ID NO: 42:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 4451 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: double  
TOPOLOGY: linear  
MOLECULE TYPE: cDNA  
US-08-717-294-42

Query Match 84.7%; Score 14.4; DB 3; Length 4451;  
Best Local Similarity 93.8%; Pred. No. 1.3e+02;  
Matches 15; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 2 GGAGCCCCAGCAATGC 17  
|||||  
4051 GGAGCCCCAGCAAGC 4066

RESULT 11  
US-09-453-702B-162  
Sequence 162, Application US/09453702B  
Patent No. 6365723  
GENERAL INFORMATION:  
APPLICANT: Blattner, Frederick R.  
Burland, Valerie  
Perna, Nicole T.  
Plunkett, Guy  
Welch, Rod  
TITLE OF INVENTION: No. 6365723el Sequences of E. coli O157  
NUMBER OF SEQUENCES: 265  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Quarles & Brady  
STREET: 1 South Pinckney Street  
CITY: Madison  
STATE: WI  
COUNTRY: US  
ZIP: 53701-2113  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Diskette, 3.50 inch, 1.44Mb storage  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: Word Perfect 8.0  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/09/453,702B  
FILING DATE: 03-Dec-1999  
CLASSIFICATION: <Unknown>  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: 60/110,955  
FILING DATE: 04-DEC-1998  
ATTORNEY/AGENT INFORMATION:  
NAME: Seay, Nicholas J.  
REGISTRATION NUMBER: 27386  
REFERENCE/DOCKET NUMBER: 960296.95017  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: (608) 251-5000  
TELEFAX: (608) 251-9166  
INFORMATION FOR SEQ ID NO: 162:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 9179  
TYPE: nucleic acid  
STRANDEDNESS: double  
TOPOLOGY: linear  
MOLECULE TYPE: DNA (genomic)  
SEQUENCE DESCRIPTION: SEQ ID NO: 162:  
US-09-453-702B-162

Query Match 84.7%; Score 14.4; DB 4; Length 9179;  
Best Local Similarity 93.8%; Pred. No. 1.4e+02;  
Matches 15; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 2 GGAGCCCCAGCAATGC 17  
|||||  
Db 2750 GCAGCCCCAGCAATGC 2765

RESULT 12  
US-08-611-280-4/c  
Sequence 4, Application US/08611280  
Patent No. 5891666  
GENERAL INFORMATION:  
APPLICANT: Matsuyama, Toshifumi  
APPLICANT: Grossman, Alex  
TITLE OF INVENTION: NOVEL GENES ENCODING LSIRF POLYPEPTIDES  
NUMBER OF SEQUENCES: 25  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Amgen Canada Inc.  
STREET: 6733 Mississauga Road, Suite 303  
CITY: Mississauga



GenCore version 5.1.3  
Copyright (c) 1993 - 2003 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: January 21, 2003, 14:54:52 ; Search time 142.611 Seconds  
(without alignments)  
268.450 Million cell updates/sec

Title: us-09-853-688-35

Perfect score: 17  
Sequence: 1 gggagcccaagcaatgc 17

Scoring table: IDENTITY\_NUC  
Gapop 10.0 , Gapext 1.0

Searched: 2185239 seqs, 1125999159 residues

Total number of hits satisfying chosen parameters: 4370478

Minimum DB seq length: 0  
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%  
Maximum Match 100%  
Listing first 45 summaries

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2: /SID52/gcgdata/geneseq/geneseq-emb1/NA1981.DAT : \*  
3: /SID52/gcgdata/geneseq/geneseq-emb1/NA1982.DAT : \*  
4: /SID52/gcgdata/geneseq/geneseq-emb1/NA1983.DAT : \*  
5: /SID52/gcgdata/geneseq/geneseq-emb1/NA1984.DAT : \*  
6: /SID52/gcgdata/geneseq/geneseq-emb1/NA1985.DAT : \*  
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8: /SID52/gcgdata/geneseq/geneseq-emb1/NA1987.DAT : \*  
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21: /SID52/gcgdata/geneseq/geneseq-emb1/NA2000.DAT : \*  
22: /SID52/gcgdata/geneseq/geneseq-emb1/NA2001A.DAT : \*  
23: /SID52/gcgdata/geneseq/geneseq-emb1/NA2001B.DAT : \*  
24: /SID52/gcgdata/geneseq/geneseq-emb1/NA2002.DAT : \*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	17	100.0	17	24	AA18882
2	17	100.0	3700	24	AA18886
3	15.4	90.6	354	21	AA18889
4	15.4	90.6	468	22	AA18891
5	15.4	90.6	468	22	AA18892
6	15.4	90.6	468	22	AA18893
7	15.4	90.6	468	22	AA18894
8	15.4	90.6	468	22	AA18895
9	15.4	90.6	468	22	AA18896

10	15.4	90.6	458	24	ABS06062	Human genome-deriv
11	15.4	90.6	710	24	ABS05786	Human ovarian anti
12	15.4	90.6	711	20	AA15243	Human gene express
13	15.4	90.6	711	20	AA198799	Human validated ca
14	15.4	90.6	785	22	AA194530	Human neuroblastom
15	15.4	90.6	805	18	AA187832	Human collagen I a2
16	15.4	90.6	852	18	AA180192	Human collagen I a2
17	15.4	90.6	1746	23	AA184464	DNA encoding novel
18	15.4	90.6	2383	22	AA194277	Human full-length
19	15.4	90.6	2820	16	AA151456	go.trka fusion use
20	15.4	90.6	3120	21	AA12510	cDNA encoding a hu
21	15.4	90.6	3409	23	AA129795	Drosophila melanog
22	15.4	90.6	4556	23	AA170098	DNA encoding novel
23	15.4	90.6	5086	24	ABN97452	Gene #3950 used to
24	15.4	90.6	5086	24	ABK64850	Human benign prost
25	15.4	90.6	5086	24	ABL92095	Human Tumour Endot
26	15.4	90.6	5086	24	ABL92105	Human Tumour Endot
27	15.4	90.6	5086	24	ABL92121	Human Tumour Endot
28	15.4	90.6	5086	24	ABL62185	Colon adenocarcino
29	15.4	90.6	5086	24	ABL62730	Colon adenocarcino
30	15.4	90.6	5145	21	AA198196	Human colon cancer
31	15.4	90.6	5416	22	ABA83119	Collagen type I al
32	15.4	90.6	5416	24	ABN95596	Gene #2094 used to
33	15.4	90.6	5416	24	ABK64492	Human benign prost
34	15.4	90.6	5416	24	ABL62092	Colon adenocarcino
35	15.4	90.6	5416	24	ABL62655	Colon adenocarcino
36	15.4	90.6	5416	24	ABL65476	Lung cancer relate
37	15.4	90.6	5564	23	ABV24693	Human prostate exp
38	15.4	90.6	8778	23	ABL29794	Drosophila melanog
39	15.4	90.6	19628	22	AA171839	Human immune/haema
40	15.4	90.6	25576	22	AA169089	Human immune/haema
41	15.4	90.6	38682	22	AA121770	Human gene for col
42	15	88.2	392	21	AA130613	Human colon cancer
43	15	88.2	1041	22	AA168387	C glutamicum codin
44	15	88.2	1087	21	AA121828	Human breast and o
45	15	88.2	1089	22	AA132204	Human DNA repair a

ALIGNMENTS

RESULT 1  
AA18882  
ID AAS18882 standard; DNA; 17 BP.  
XX  
AC AAS18882;  
XX  
DT 12-MAR-2002 (first entry)  
XX  
DE Growth hormone 1 gene (GH1) specific fragment, PCR primer GH1F.  
XX  
KW Growth hormone 1; GH1; osteopathic; gene therapy; protein therapy;  
KW diabetes; obesity; infection; acromegaly; gigantism; sodium retention;  
KW water retention; metabolic syndrome; mood disorder; sleep disorder;  
KW Growth hormone dysfunction; familial growth hormone deficiency;  
KW short stature; pituitary storage defect; human; PCR primer; GH1F; ss.  
XX Homo sapiens.  
XX WO200185993-A2.  
XX  
PD 15-NOV-2001.  
XX  
PF 14-MAY-2001; 2001WO-GB02126.  
XX  
PR 12-MAY-2000; 2000GB-0011459.  
PR 14-JUL-2000; 2000EP-0306004.  
XX  
FA (UYWA-) UNIV WALES COLLEGE OF MEDICINE.  
XX  
PI Cooper DN, Procter AM, Gregory J, Millar DS;  
XX Probe #5562 for ge  
XX WPI; 2002-089798/12.  
DR

```

XX Detecting growth hormone variants (GHI), useful in screening patients
PT for growth hormone irregularities, comprises comparing the nucleotide
PT sequence of a GHI gene from a test sample with that of a standard
PT sequence of the human GHI
XX
XX Example 2; Page 39; 95pp; English.
XX
XX The invention described a method of detecting variation in growth hormone
CC l (GHI), and therefore GH dysfunction in an individual. The method
CC comprises comparing the nucleotide sequence of GHI gene obtained from the
CC test sample with a standard human GHI gene sequence, in order to identify
CC variation (GHI variant). The method is useful in screening patients for
CC growth hormone irregularities or producing variant proteins for treating
CC irregularities, and for the early detection and appropriate clinical
CC management of familial GH deficiency. The GHI variants are useful in
CC therapeutic, diagnostic or detection method, particularly for determining
CC binding defects and susceptibility to a disease such as diabetes, obesity
CC or infection; for treating acromegaly or gigantism conditions associated
CC with lactogenic, diabetogenic, lipolytic and protein anabolic effects,
CC conditions associated with sodium and water retention, metabolic
CC syndromes, mood and sleep disorders; diagnosing GH dysfunction and
CC determining pituitary storage defects. The GHI variants are especially
CC useful in gene therapy or protein therapy. The GHI or GH variant may also
CC be used in the preparation of a medicament, diagnostics composition or
CC kit, or detection kit. The method has the advantage of: expanding the
CC know spectrum of GHI gene mutations; evaluating the role of GHI gene
CC mutations in the etiology of short stature; identifying of the mode of
CC inheritance of novel lesions; evaluation the effects of GHI mutations on
CC the structure and function of the GH molecule and development of rapid
CC diagnostic tests for inherited GH deficiency. This sequence is the GHI
CC PCR primer. GHIF, used with GHI1R (AAS18883) to amplify a GHI-specific
CC fragment, described in the method of the invention.
XX
XX Sequence 17 BP; 4 A; 6 C; 6 G; 1 T; 0 other;
SQ
Query Match 100.0%; Score 17; DB 24; Length 17;
Best Local Similarity 100.0%; Pred. No. 46;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGGAGCCCGAGCAATGC 17
Db 1 GGGAGCCCGAGCAATGC 17

RESULT 2
AAS18886
ID AAS18886 standard; DNA; 3700 BP.
XX
XX AAS18886;
AC
XX
D1 12-MAR-2002 (first entry)
D2
D3
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FT		/tag= ab	"Single nucleotide polymorphism"
FT	variation	/standard_name= replace(573,G)	"Single nucleotide polymorphism"
FT		/tag= ac	
FT		/standard_name= replace(580,G)	"Single nucleotide polymorphism"
FT	variation	/tag= ad	"Single nucleotide polymorphism"
FT		/standard_name= replace(585,G)	
FT	variation	/tag= ae	"Single nucleotide polymorphism"
FT		/standard_name= replace(586,T)	"Single nucleotide polymorphism"
FT	variation	/tag= af	
FT		/standard_name= replace(620,A)	"Single nucleotide polymorphism"
FT	variation	/tag= ag	"Single nucleotide polymorphism"
FT		/standard_name= replace(622,G)	
FT	variation	/tag= ah	"Single nucleotide polymorphism"
FT		/standard_name= replace(649,C)	"Single nucleotide polymorphism"
FT	variation	/tag= ai	"Single nucleotide polymorphism"
FT		/standard_name= replace(665,C)	
FT	variation	/tag= aj	"Single nucleotide polymorphism"
FT		/standard_name= replace(670,G)	
FT	variation	/tag= ak	"Single nucleotide polymorphism"
FT		/standard_name= replace(676,A)	"Single nucleotide polymorphism"
FT	variation	/tag= al	"Single nucleotide polymorphism"
FT		/standard_name= replace(685,A)	
FT	variation	/tag= am	"Single nucleotide polymorphism"
FT	exon	/standard_name= 701..772	
FT		/tag= an	
FT	intron	/number= 1	
FT		773..1032	
FT		/tag= ao	
FT		/number= 1	
FT	variation	replace(836,C)	
FT		/tag= ap	"Single nucleotide polymorphism"
FT	variation	/standard_name= replace(839,C)	
FT		/tag= aq	
FT	variation	/standard_name= replace(879,G)	"Single nucleotide polymorphism"
FT		/tag= ar	
FT	variation	/standard_name= replace(883,A)	"Single nucleotide polymorphism"
FT		/tag= as	
FT	variation	/standard_name= replace(901,C)	"Single nucleotide polymorphism"
FT		/tag= at	
FT	variation	/standard_name= replace(1010,T)	"Single nucleotide polymorphism"
FT		/tag= au	
FT	exon	/standard_name= 1033..1193	"Single nucleotide polymorphism"
FT		/tag= av	
FT		/number= 2	
FT	variation	replace(1097,A)	
FT		/tag= aw	"Single nucleotide polymorphism"
FT	variation	/standard_name= replace(1101,T)	
FT		/tag= ax	
FT	mat_peptide	/standard_name= 1101..2227	"Single nucleotide polymorphism"
FT		/tag= ay	
FT	variation	/label= "mature_GH1"	
FT		replace(1114,T)	
FT		/tag= az	

FT	variation	/standard_name= replace(1169,A) /tag= ba	"Single nucleotide polymorphism"
FT			
FT	variation	/standard_name= replace(1182,T) /tag= bb	"Single nucleotide polymorphism"
FT			
FT	variation	/standard_name= replace(1189,G) /tag= bc	"Single nucleotide polymorphism"
FT			
FT	variation	/standard_name= replace(1193,G) /tag= bd	"Single nucleotide polymorphism"
FT			
FT	intron	/standard_name= 1194..1402 /tag= be	"Single nucleotide polymorphism"
FT		/number= 2	
FT	variation	replace(1196,G) /tag= bf	"Single nucleotide polymorphism"
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FT	variation	/standard_name= replace(1196,C) /tag= bq	"Single nucleotide polymorphism"
FT			
FT	variation	/standard_name= replace(1208,C) /tag= bh	"Single nucleotide polymorphism"
FT			
FT	variation	/standard_name= replace(1212,T) /tag= bi	"Single nucleotide polymorphism"
FT			
FT		/standard name=	"Single nucleotide polymorphism"

RESULT 3  
AAC30859/C  
ID AAC30859 standard; cDNA; 354 BP.  
XX  
XX  
AAC30859;  
AC  
XX  
XX  
06-OCT-2000 (first entry)  
DT  
XX  
XX  
XX  
DE Human secreted protein 5' EST, SEQ ID NO: 34934.

FT mat\_peptide

```
FT      /*tag= ay
FT      /label= "mature_CH1"
~m      variation
        replace(114,T)
        /*tag= az
```

CC The present sequence is one of a large number of 5' ESTs derived from  
CC mRNAs encoding secreted proteins. No ORF has yet been conclusively  
CC identified within the present sequence. The 5' ESTs were prepared from  
CC total human RNAs or polyA+ RNAs derived from 30 different tissues. EST  
CC sequences usually correspond mainly to the 3' untranslated region (UTR)  
CC of the mRNA because they are often obtained from oligo-dT primed cDNA  
CC libraries. Such ESTs are not well suited for isolating cDNA sequences  
CC derived from the 5' ends of mRNAs and even in those cases where longer  
CC cDNA sequences have been obtained, the full 5' UTR is rarely included.  
CC 5' ESTs are derived from mRNAs with intact 5' ends and can therefore be  
CC used to obtain full length cDNAs and genomic DNAs. 5' ESTs are also used  
CC in diagnostic, forensic, gene therapy and chromosome mapping procedures.  
CC They are used to obtain upstream regulatory sequences and to design  
CC expression and secretion vectors.

XX  
SQ Sequence 354 BP; 90 A; 89 C; 79 G; 96 T; 0 other;

Query Match 90.6%; Score 15.4; DB 21; Length 354;  
Best Local Similarity 94.1%; Pred. No. 3.1e+02;  
Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 GGGAGCCCCAGCAATGC 17  
|||||||  
Db 148 GGGAGCCCCAGCAATGC 132

## RESULT 4

ABA57641  
ID ABA57641 standard; DNA: 468 BP.

XX  
AC ABA57641;

XX  
DT 01-FEB-2002 (first entry)

XX  
DE Human foetal liver: single exon nucleic acid probe #5946.

XX  
KW Human; foetal liver; gene expression; single exon nucleic acid probe; ss.

XX  
OS Homo sapiens.

XX  
PN W0200157277-A2.

XX  
PD 09-AUG-2001.

XX  
PF 30-JAN-2001; 2001WO-US00669.

XX  
PR 04-FEB-2000; 2000US-0180312.

XX  
PR 26-MAY-2000; 2000US-0207456.

XX  
PR 30-JUN-2000; 2000US-0608408.

XX  
PR 03-AUG-2000; 2000US-0632366.

XX  
PR 21-SEP-2000; 2000US-0234687.

XX  
PR 27-SEP-2000; 2000US-0236359.

XX  
PR 04-OCT-2000; 2000GB-0024263.

XX  
PA (MOLE-) MOLECULAR DYNAMICS INC.

XX  
PI Penn SG, Hanzel DK, Chen W, Rank DR;

XX  
XX WPI; 2001-483447/52.

XX  
XX Human genome-derived single exon nucleic acid probes useful for  
XX analyzing gene expression in human fetal liver -

XX  
XX Claim 1; SEQ ID NO 5946; 639pp + sequence listing; English.

XX  
CC The invention relates to a single exon nucleic acid probe for  
CC measuring human gene expression in a sample derived from human foetal

CC liver. The single exon nucleic acid probes may be used for predicting,  
CC measuring and displaying gene expression in samples derived from human

CC foetal liver. The present sequence is a single exon nucleic acid  
CC probe of the invention.

CC  
CC Note: The sequence data for this patent did not form part of the  
GC printed specification, but was obtained in electronic format directly

CC from WIPO at ftp.wipo.int/pub/published\_pct\_sequences.

XX  
SQ Sequence 468 BP; 113 A; 116 C; 146 G; 93 T; 0 other;

Query Match 90.6%; Score 15.4; DB 22; Length 468;  
Best Local Similarity 94.1%; Pred. No. 3.1e+02;

Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 GGGAGCCCCAGCAATGC 17  
|||||||  
Db 416 GGGAGCCCCAGCAATTC 432

## RESULT 5

ABA27062  
ID ABA27062 standard; DNA: 468 BP.

XX  
AC ABA27062;

XX  
DT 23-JAN-2002 (first entry)

XX  
DE Probe #5528 for gene expression analysis in human heart cell sample.

XX  
KW Human; gene expression; heart; microarray; vascular system; probe;

XX  
KW cardiovascular disease; hypertension; cardiac arrhythmia;

XX  
KW congenital heart disease; ss.

XX  
OS Homo sapiens.

XX  
PN W0200157274-A2.

XX  
PD 09-AUG-2001.

XX  
PF 30-JAN-2001; 2001WO-US00666.

XX  
PR 04-FEB-2000; 2000US-0180312.

XX  
PR 26-MAY-2000; 2000US-0207456.

XX  
PR 30-JUN-2000; 2000US-0608408.

XX  
PR 03-AUG-2000; 2000US-0632366.

XX  
PR 21-SEP-2000; 2000US-0234687.

XX  
PR 27-SEP-2000; 2000US-0236359.

XX  
PR 04-OCT-2000; 2000GB-0024263.

XX  
PA (MOLE-) MOLECULAR DYNAMICS INC.

XX  
PI Penn SG, Hanzel DK, Chen W, Rank DR;

XX  
XX WPI; 2001-488899/53.

XX  
XX Single exon nucleic acid probes for analyzing gene expression in human

XX  
XX hearts -

XX  
PS Claim 1; SEQ ID NO 5528; 530pp; English.

XX  
CC The present invention relates to single exon nucleic acid probes for

CC measuring human gene expression in a sample derived from human heart. The

CC present sequence is one such probe. The probes may be used for

CC predicting, measuring and displaying gene expression in samples derived

CC from the human heart via microarrays. By measuring gene expression, the

CC probes are useful for predicting, diagnosing, grading, staging,

CC monitoring and prognosing diseases of the human heart and vascular system

CC e.g. cardiovascular disease, hypertension, cardiac arrhythmias and

CC congenital heart disease.

CC Note: The sequence data for this patent did not form part of the printed

CC specification, but was obtained in electronic format directly from WIPO

CC at ftp.wipo.int/pub/published\_pct\_sequences.

XX  
SQ Sequence 468 BP; 113 A; 116 C; 146 G; 93 T; 0 other;

Query Match 90.6%; Score 15.4; DB 22; Length 468;  
Best Local Similarity 94.1%; Pred. No. 3.1e+02;  
Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

```
QY 1 GGGAGCCCCCAGCAATGC 17
Db 416 GGGAGCCCCCAGCAATTC 432

RESULT 6
AAK05695
ID AAK05695 standard; DNA; 468 BP.
AC AAK05695;
XX
XX
XX 05-NOV-2001 (first entry)
DT
DE human brain expressed single exon probe SEQ ID NO: 5686.
XX
XX human; brain expressed exon; gene expression analysis; probe;
KW microarray; Alzheimer's disease; multiple sclerosis; schizophrenia;
KW epilepsy; cancer; ss.
XX
XX Homo sapiens.
OS
XX
XX WO200157275-A2.
PN
XX
XX 09-AUG-2001.
PD
XX
XX 30-JAN-2001; 2001WO-US00667.
PF
XX
XX 04-FEB-2000; 2000US-0180312.
PR
XX 26-MAY-2000; 2000US-0207456.
PR
XX 30-JUN-2000; 2000US-0608408.
PR
XX 03-AUG-2000; 2000US-0632366.
PR
XX 21-SEP-2000; 2000US-0234687.
PR
XX 27-SEP-2000; 2000US-0236359.
PR
XX 04-OCT-2000; 2000GB-0024263.
XX
XX (MOLE-) MOLECULAR DYNAMICS INC.
PA
XX
XX Penn SG, Hanzel DK, Chen W, Rank DR;
PI
XX
XX WPI; 2001-488900/53.
XX
XX Human genome-derived single exon nucleic acid probes useful for
PT analyzing gene expression in human bone marrow.
PT
XX
XX Example 4; SEQ ID NO: 5687; 658pp + Sequence Listing; English..
PS
XX
XX The present invention provides a number of single exon nucleic acid
CC probes which are derived from genomic sequences expressed in the human
CC bone marrow. They can be used to measure gene expression in bone marrow
CC samples, which may enable the improved diagnosis and treatment of cancers
CC such as lymphoma, leukaemia and myeloma. The present sequence is one of
CC the probes of the invention.
XX
XX
XX Sequence 468 BP; 113 A; 116 C; 146 G; 93 T; 0 other;
SQ

Query Match 90.6%; Score 15.4; DB 22; Length 468;
Best Local Similarity 94.1%; Pred. NO. 3.1e+02;
Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 GGGAGCCCCCAGCAATGC 17
Db 416 GGGAGCCCCCAGCAATTC 432

RESULT 8
AAI15629
ID AAI15629 standard; DNA; 468 BP.
XX
XX
XX AAI15629;
AC
XX
XX
XX 12-OCT-2001 (first entry)
DT
XX
XX
XX Probe #5562 for gene expression analysis in human cervical cell sample.
DE
XX
XX Probe; human; microarray; gene expression; cervical epithelial cell;
KW cervical cancer; ss.
XX
XX
XX Homo sapiens.
OS
XX
XX WO200157278-A2.
PN
XX
XX 09-AUG-2001.
PD
XX
XX 30-JAN-2001; 2001WO-US00670.
PF
XX
XX 04-FEB-2000; 2000US-0180312.
PR
XX 26-MAY-2000; 2000US-0207456.
PR
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```
QY 1 GGGAGCCCCCAGCAATGC 17
Db 416 GGGAGCCCCCAGCAATTC 432

RESULT 7
AAK31310
ID AAK31310 standard; DNA; 468 BP.
AC AAK31310;
XX
XX 06-NOV-2001 (first entry)
DT
DE human brain expressed single exon probe SEQ ID NO: 5686.
XX
XX human; brain expressed exon; gene expression analysis; probe;
KW microarray; Alzheimer's disease; multiple sclerosis; schizophrenia;
KW epilepsy; cancer; ss.
XX
XX Homo sapiens.
OS
XX
XX WO200157275-A2.
PN
XX
XX 09-AUG-2001.
PD
XX
XX 30-JAN-2001; 2001WO-US00667.
PF
XX
XX 04-FEB-2000; 2000US-0180312.
PR
XX 26-MAY-2000; 2000US-0207456.
PR
XX 30-JUN-2000; 2000US-0608408.
PR
XX 03-AUG-2000; 2000US-0632366.
PR
XX 21-SEP-2000; 2000US-0234687.
PR
XX 27-SEP-2000; 2000US-0236359.
PR
XX 04-OCT-2000; 2000GB-0024263.
XX
XX (MOLE-) MOLECULAR DYNAMICS INC.
PA
XX
XX Penn SG, Hanzel DK, Chen W, Rank DR;
PI
XX
XX WPI; 2001-483446/52.
XX
XX Single exon nucleic acid probes for analyzing gene expression in human
PT brains.
PT
XX
XX Example 4; SEQ ID NO: 5686; 650pp + Sequence Listing; English.
PS
XX
XX The present invention provides a number of single exon nucleic acid
CC probes which are derived from genomic sequences expressed in the human
CC brain. They can be used to measure gene expression in brain cell samples,
CC which may enable the diagnosis and improved treatment of nervous system
CC diseases such as Alzheimer's disease, multiple sclerosis, schizophrenia,
CC epilepsy and cancers. The present sequence is one of the probes of the
CC invention.
XX
XX
XX Sequence 468 BP; 113 A; 116 C; 146 G; 93 T; 0 other;
SQ

Query Match 90.6%; Score 15.4; DB 22; Length 468;
Best Local Similarity 94.1%; Pred. NO. 3.1e+02;
Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 GGGAGCCCCCAGCAATGC 17
Db 416 GGGAGCCCCCAGCAATTC 432

RESULT 7
AAK31310
ID AAK31310 standard; DNA; 468 BP.
AC AAK31310;
XX
XX 06-NOV-2001 (first entry)
DT
DE human brain expressed single exon probe SEQ ID NO: 5686.
XX
XX human; brain expressed exon; gene expression analysis; probe;
KW microarray; Alzheimer's disease; multiple sclerosis; schizophrenia;
KW epilepsy; cancer; ss.
XX
XX Homo sapiens.
OS
XX
XX WO200157275-A2.
PN
XX
XX 09-AUG-2001.
PD
XX
XX 30-JAN-2001; 2001WO-US00667.
PF
XX
XX 04-FEB-2000; 2000US-0180312.
PR
XX 26-MAY-2000; 2000US-0207456.
PR
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PR 30-JUN-2000; 2000US-0608408.  
PR 03-AUG-2000; 2000US-0532366.  
PR 21-SEP-2000; 2000US-0234687.  
PR 27-SEP-2000; 2000US-0236359.  
PR 04-OCT-2000; 2000GB-0024263.  
XX  
XX (MOLE-) MOLECULAR DYNAMICS INC.  
XX  
XX Penn SG, Hanzel DK, Chen W, Rank DK;  
XX WPI; 2001-488901/53.  
XX  
XX Human genome-derived single exon nucleic acid probes useful for  
PT analyzing gene expression in human cervical epithelial cells -  
XX  
XX Claim 25; SEQ ID NO 5562; 487pp; English.  
XX  
XX The present invention relates to human single exon nucleic acid probes  
CC (SENP). The present sequence is one such probe. The SENPs are derived  
CC from human HeLa cells. The SENPs can be used to produce a single exon  
CC microarray, which can be used for measuring human gene expression in a  
CC sample derived from human cervical epithelial cells. By measuring gene  
CC expression, the probes are therefore useful in grading and/or staging  
CC of diseases of the cervix, notably cervical cancer.  
CC Note: The sequence data for this patent did not form part of the printed  
CC specification, but was obtained in electronic format directly from WIPO  
CC at ftp.wipo.int/pub/published\_pct\_sequences.  
XX  
XX Sequence 468 BP; 113 A; 116 C; 146 G; 93 T; 0 other;  
SQ  
Query Match 90.6%; Score 15.4; DB 22; Length 468;  
Best Local Similarity 94.1%; Pred. No. 3.1e+02;  
Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;  
OY 1 GGGAGCCCCAGCAATGC 17  
Db 416 GGGAGCCCCAGCAATTC 432  
RESULT 9  
AA137207  
ID AA137207 standard; DNA; 468 BP.  
AC AA137207;  
XX  
XX 17-OCT-2001 (first entry)  
XX  
XX Probe #5893 used to measure gene expression in human placenta sample.  
DE  
XX Probe; microarray; human; placenta; antenatal diagnosis;  
KW genetic disorder; ss.  
XX  
XX Homo sapiens.  
OS  
XX W0200157272-A2.  
XX  
XX 09-AUG-2001.  
XX  
XX 30-JAN-2001; 2001WO-US00663.  
XX  
XX 04-FEB-2000; 2000US-0180312.  
PR 26-MAY-2000; 2000US-0207456.  
PR 30-JUN-2000; 2000US-0608408.  
PR 03-AUG-2000; 2000US-0632366.  
PR 21-SEP-2000; 2000US-0234687.  
PR 27-SEP-2000; 2000US-0236359.  
PR 04-OCT-2000; 2000GB-0024263.  
XX  
XX (MOLE-) MOLECULAR DYNAMICS INC.  
XX  
XX Penn SG, Hanzel DK, Chen W, Rank DK;  
XX WPI; 2001-488901/53.  
XX

XX Human genome-derived single exon nucleic acid probes useful for  
PT analyzing gene expression in human placenta -  
XX  
XX Claim 25; SEQ ID NO 5893; 654pp; English.  
XX  
XX The present invention relates to single exon nucleic acid probes (SENP).  
CC The present sequence is one such probe. The probes are useful for  
CC producing a microarray for predicting, measuring and displaying gene  
CC expression in samples derived from human placenta. The probes are useful  
CC for antenatal diagnosis of human genetic disorders.  
XX  
XX Sequence 468 BP; 113 A; 116 C; 146 G; 93 T; 0 other;  
SQ  
Query Match 90.6%; Score 15.4; DB 22; Length 468;  
Best Local Similarity 94.1%; Pred. No. 3.1e+02;  
Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;  
OY 1 GGGAGCCCCAGCAATGC 17  
Db 416 GGGAGCCCCAGCAATTC 432  
RESULT 10  
ABS06062  
ID ABS06062 standard; DNA; 468 BP.  
XX  
XX ABS06062;  
XX  
XX 19-AUG-2002 (first entry)  
XX  
XX Human genome-derived single exon probe from lung SEQ ID NO 6053.  
DE  
XX Human; ds; single exon probe; asthma; lung cancer; COPD; ILD;  
KW chronic obstructive pulmonary disease; interstitial lung disease;  
KW familial idiopathic pulmonary fibrosis; neurofibromatosis;  
KW tuberous sclerosis; Gaucher's disease; Niemann-Pick disease;  
KW hermannsky-pudlak syndrome; sarcoidosis; pulmonary haemorrhoidosis;  
KW pulmonary histiocytosis; lymphangioleiomyomatosis; karagener syndrome;  
KW pulmonary alveolar proteinosis; fibrocystic pulmonary dysplasia;  
KW primary ciliary dyskinesia; pulmonary hypertension;  
KW hyaline membrane disease.  
XX  
XX Homo sapiens.  
OS  
XX W0200186003-A2.  
XX  
XX 15-NOV-2001.  
XX  
XX 30-JAN-2001; 2001WO-US00665.  
XX  
XX 04-FEB-2000; 2000US-180312P.  
PR 26-MAY-2000; 2000US-207456P.  
PR 30-JUN-2000; 2000US-0608408.  
PR 03-AUG-2000; 2000US-0632366.  
PR 21-SEP-2000; 2000US-234687P.  
PR 27-SEP-2000; 2000US-236359P.  
PR 04-OCT-2000; 2000GB-0024263.  
XX  
XX (MOLE-) MOLECULAR DYNAMICS INC.  
XX  
XX Penn SG, Hanzel DK, Chen W, Rank DK;  
XX WPI; 2002-114183/15.  
XX  
XX Spatially-addressable set of single exon nucleic acid probes, used to  
PT measure gene expression in human lung samples -  
XX  
XX Claim 1; SEQ ID NO 6053; 634pp; English.  
XX  
XX The invention relates to a spatially-addressable set of single exon  
CC nucleic acid probes for measuring gene expression in a sample derived  
CC from human lung comprising single exon nucleic acid probes having one of

12614 nucleic acid sequences mentioned in the specification, or their complements or the 12387 open reading frames derived from the 12614 probes. Also included are a microarray comprising the novel set of probes; the novel set of probes which hybridise at high stringency to a nucleic acid expressed in the human lung; measuring gene expression in a sample derived from human lung, comprising (a) contacting the array with a collection of detectably labeled nucleic acids derived from human lung mRNA, and (b) measuring the label detectably bound to each probe of the array; identifying exons in a eukaryotic genome, comprising (a) algorithmically predicting at least one exon from genomic sequences of the eukaryote; and (b) detecting specific hybridisation of detectably labeled nucleic acids from eukaryote lung mRNA, to a single exon probe. In the above mentioned microarray; assigning exons to a single gene, comprising (a) identifying exons from genomic sequence by the method above and (b) measuring the expression of each of the exons in several tissues and/or cell types using hybridisation to a single exon microarrays having a probe with the exon, where a common pattern of expression of the exons in the tissues and/or cell types indicates that the exons should be assigned to a single gene; a peptide comprising one of 12011 sequences, mentioned in the specification, or encoded by the probes/open reading frames (ORF). The probes are used for gene expression analysis, and for identifying exons in a gene, particularly using human lung derived mRNA and for the study of lung diseases such as asthma, lung cancer, chronic obstructive pulmonary disease (COPD), interstitial lung disease (ILD), familial idiopathic pulmonary fibrosis, neurofibromatosis, tuberculous sclerosis, Gaucher's disease, Niemann-Pick disease, Hermansky-Pudlak syndrome, sarcoidosis, pulmonary haemoderiosis, pulmonary histiocytosis, lymphangioleiomyomatosis, pulmonary alveolar proteinosis, Karagener syndrome, fibrocystic pulmonary dysplasia, primary ciliary dyskinesia, pulmonary hypertension and hyaline membrane disease. The present sequence is a single exon probe of the invention.

Note: The sequence data for this patent did not form part of the printed specification, but was obtained in electronic format directly from WIPO at [ftp.wipo.int/pub/published\\_pct\\_sequences](http://ftp.wipo.int/pub/published_pct_sequences).

XX SQ Sequence 468 BP; 113 A; 116 G; 146 G; 93 T; 0 other;

Query Match 90.6%; Score 15.4; DB 24; Length 468;  
Best Local Similarity 94.1%; Pred. No. 3.1e+02;  
Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 GGGAGCCCCAGCAATGC 17  
|||||  
DB 416 GGGAGCCCCAGCAATTC 432

RESULT 11  
ABQ55786  
ID ABQ55786 standard; cDNA; 710 BP.

XX AC ABQ55786;

DT 22-AUG-2002 (first entry)

DE Human ovarian antigen HOPKO61 cDNA, SEQ ID NO:1666.

XX KW Human; ovarian antigen; ovary; ovarian; breast; cancer; tumour;  
KW ovarian cancer; breast cancer; tumour; reproductive system disorder;  
KW infertility; pregnancy disorder; anovulation; polycystic ovary syndrome;  
KW PCOS; ovarian cyst; dysmenorrhea; endocrine disorder; infection;  
KW inflammatory condition; immune disorder; blood disorder;  
KW cardiovascular disorder; respiratory disorder; neurological disorder;  
KW gastrointestinal disorder; urinary system disorder; drug screening;  
KW gene therapy; chromosome mapping; forensic analysis;  
KW antibody preparation; cytostatic; immunomodulatory; neuroprotective;  
KW antiinflammatory; gynaecological; reproductive; gene; ss.

OS Homo sapiens.

XX WO200200677-A1.

Human gene expression product cDNA sequence SEQ ID NO:2712.

XX 03-JAN-2002.  
PD  
XX 07-JUN-2001; 2001WO-US18569.  
PF  
XX 07-JUN-2000; 2000US-209467P.  
PR  
XX (HUMA-) HUMAN GENOME SCI INC.  
PA

PI Birse CE, Rosen CA;  
XX  
DR WPI: 2002-147878/19.  
DR P-PSDB; ARP42709.

XX Isolated nucleic acid molecules encoding novel ovarian polypeptides,  
PT useful in the prevention, treatment and diagnosis of cancer (e.g.  
PT ovarian cancer), immune disorders, cardiovascular disorders and  
PT neurological diseases -

XX Claim 1: SEQ ID No 1666; 2922pp; English.

XX The invention relates to 2175 novel human ovarian antigens (ABP41054-  
CC ABP43228) and to cDNAs encoding them (ABQ54131-ABQ56305), and also  
CC encompasses polypeptides 90% identical and polynucleotides 95% identical  
CC to the sequences of the invention. The invention additionally relates to  
CC recombinant vectors and host cells comprising human ovarian antigen  
CC polynucleotides, antibodies against human ovarian antigens, and the use  
CC of ovarian antigen polynucleotides and polypeptides in diagnosing,  
CC treating, prognosing or preventing various ovary and/or breast-related  
CC disorders. Such conditions include ovarian cancer and breast cancer, and  
CC metastatic tumours of ovarian or breast origin, reproductive system  
CC disorders (e.g., infertility, disorders of pregnancy, anovulation,  
CC polycystic ovary syndrome, ovarian cysts, and dysmenorrhea), endocrine  
CC disorders, infections (e.g., chlamydia, HIV, toxoplasmosis, and toxic  
CC shock syndrome), inflammatory conditions (e.g., mastitis, oophoritis and  
CC vaginitis), immune disorders (e.g., congenital and acquired  
CC immunodeficiencies, autoimmune oophoritis, systemic lupus erythematosus),  
CC blood-related disorders (e.g., anaemia), cardiovascular disorders,  
CC respiratory disorders, neurological disorders, gastrointestinal disorders  
CC and urinary system disorders. Ovarian antigen polypeptides and  
CC polynucleotides may also be used in screening for compounds which  
CC modulate ovarian antigen expression or activity. The polynucleotides may  
CC further be used for gene therapy, chromosome mapping, in the  
CC identification of individuals and in forensic analysis, and the  
CC polypeptides may be used as food additives or to prepare antibodies  
CC useful in disease diagnosis, drug targeting and phenotyping. The present  
CC sequence represents cDNA encoding a human ovarian antigen of the  
CC invention.

CC Note: The sequence data for this patent did not form part of the printed  
CC specification, but was obtained in electronic format directly from WIPO  
CC at [ftp.wipo.int/pub/published\\_pct\\_sequences](http://ftp.wipo.int/pub/published_pct_sequences).

XX SQ Sequence 710 BP; 113 A; 244 C; 222 G; 125 T; 6 other;

Query Match 90.6%; Score 15.4; DB 24; Length 710;  
Best Local Similarity 94.1%; Pred. No. 3.2e+02;  
Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 GGGAGCCCCAGCAATGC 17  
|||||  
DB 474 GGGAGCCCCAGCAATGC 490

RESULT 12  
AAZ15243  
ID AAZ15243 standard; cDNA; 711 BP.

XX AAZ15243;

XX 12-OCT-1999 (first entry)

Human: gene: gene expression product; diagnosis; therapy; probe; detection; mapping; tissue typing; profiling; forensic; cancer; genetic analysis; colorectal cancer; breast cancer; lung cancer; ss.

Human sapiens.

W09938972-A2.

05-AUG-1999.

28-JAN-1999; 99WO-US01619.

03-APR-1998; 98US-0080666.

28-JAN-1998; 98US-0072910.

24-FEB-1998; 98US-0075954.

31-MAR-1998; 98US-0080114.

03-APR-1998; 98US-0080515.

(CHIR ) CHIRON CORP.

(HYSE-) HYSEQ INC.

Crkvenjakov R, Drmanac R, Drmanac S;

Escobedo J, Garcia PD, Garcia V, Giese K, Innis MA;

Jones WL, Kassam A, Kennedy GC, Kita D, Labat I;

Lamson G, Leshkowitz D, Pot D, Randazzo F, Reinhard C;

Stache-Crain B, Sudduth-Klinger J, Williams LT;

WPI: 1999-494092/41.

Novel human genes and their expression products which are differentially expressed in different cell types

Claim 1: Page 1317; 2479pp; English.

The present invention describes a library of human polynucleotides comprising the sequences given in AA212532 to AA217779. Also described is a method of detecting differentially expressed genes correlated with the cancerous state of a mammalian cell, comprising detecting at least one differentially expressed gene product in a test sample from a cell suspected of being cancerous, where the gene product is encoded by one of the 5248 polynucleotide sequences given in AA212532 to AA217779. The polynucleotides can be used as a source of primers and probes, which can be used for a variety of purpose, e.g. detection of expression levels, mapping, tissue typing or profiling, forensics, genetic analysis and detection of polymorphisms. Polypeptides encoded by the polynucleotides can be used for raising antibodies for experimental, diagnostic and therapeutic purposes. The polynucleotides may also be used to construct arrays for diagnostics (which may be used to determine function of an encoded protein); and to detect differences in expression levels between two cells (e.g. to identify abnormal or diseased tissue in a human, to identify a genetic predisposition or susceptibility to a disease such as cancer). The polynucleotides of the invention are especially used in the diagnosis, prognosis and management of colorectal cancer, breast cancer, and lung cancer. The polynucleotides can also be used to screen for peptide analogues and antagonists.

Sequence 711 BP: 137 A; 214 C; 194 G; 159 T; 7 other;

Query Match 90.6%; Score 15.4; DB 20; Length 711;

Best Local Similarity 94.1%; Pred. No. 3.2e+02;

Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 GGGAGCCCCCAGCATGC 17

|||||

Db 478 GGGAGCCCCCAGCAACG 494

RESULT 13

AAx98799

ID AAx98799 standard; cDNA: 711 BP.

XX

XX AAx98799;

XX

DT 24-SEP-1999 (first entry)

XX Human validated cancer cell derived cDNA #121.

XX

XX Cancer: human; colon; breast; lung; transmembrane receptor; ATPase;

KW integral membrane protein; aspartyl protease; GATA family; wnt family;

KW transcription factor; G-protein alpha subunit; protein phosphatase;

KW phospholipase binding protein; diacylglycerol binding protein; trypsin;

KW protein kinase; tyrosine phosphatase; developmental signalling protein;

KW WW/rsp5/WMP domain; therapy; forensic; genetic mapping; diagnostic;

KW detection; treatment; cervical; melanoma; colorectal adenocarcinoma;

KW Wilm's tumour; retinoblastoma; sarcoma; myosarcoma; lung carcinoma;

KW leukemia; lymphoma; dysplasia; hyperplasia; endometrium; adrenal;

XX prostate; ss.

XX Homo sapiens.

OS

XX W09933982-A2.

PN

XX

XX 08-JUL-1999.

PD

XX

XX 22-DEC-1998; 98WO-US27610.

PF

XX

XX 21-DEC-1998; 98US-0217471.

PR

XX 23-DEC-1997; 97US-0068755.

PR

XX 03-APR-1998; 98US-0080664.

PR

XX 21-OCT-1998; 98US-0105234.

PR

XX 27-OCT-1998; 98US-0105877.

XX

XX (CHIR ) CHIRON CORP.

PA

XX (HYSE-) HYSEQ INC.

XX

XX Crkvenjakov R, Drmanac R, Drmanac S;

PI Escobedo J, Garcia PD, Garcia V, Giese K, Innis MA;

PI Jones WL, Kassam A, Kennedy GC, Kita D, Labat I;

PI Lamson G, Leshkowitz D, Pot D, Randazzo F, Reinhard C;

PI Stache-Crain B, Sudduth-Klinger J, Williams LT;

XX

DR WPI: 1999-430243/36.

XX

XX New isolated human polynucleotides

PT

XX

XX Claim 1: Page 462; 591pp; English.

PS

XX This invention describes novel isolated human polynucleotides obtained by screening for differential expression in colon cancer, breast cancer and lung cancer cell lines. The polynucleotides of the invention are represented in AAx98275-X99118 and encode polypeptides of protein families selected from 4 transmembrane segments integral membrane proteins, 7 transmembrane receptors, ATPases associated with various cellular activities (AAA), eukaryotic aspartyl proteases, GATA family of transcription factors, G-protein alpha subunit, phospholipase or diacylglycerol binding proteins, protein kinase, protein phosphatase 2C, protein tyrosine phosphatase, trypsin, wnt family of developmental signalling proteins and WW/rsp5/WMP domain containing proteins. The encoded polypeptides also have a functional domain selected from Ank repeat, basic region plus leucine zipper transcription factors, bromodomain, EF-hand, SH3 domain, WD domain/G-beta repeats, zinc finger (C2H2 type), zinc finger (CCHC class), and zinc-binding metalloprotease domain. The polynucleotides encode polypeptides with similarity to known protein families and are predicted to have similar properties. The novel polynucleotides can be used to develop products for use as therapeutic agents and in forensics, genetic analysis, mapping and diagnostic applications. In particular, the product can be used for the detection and management of cancers. They can be used for treating e.g. cervical cancers, melanomas, colorectal adenocarcinomas, Wilm's tumour, sarcomas, retinoblastoma, myosarcomas, lung carcinomas, leukemias, such as chronic myelogenous leukemia, promyelocytic leukemia, monocytic leukemia, and myeloid leukemia, and lymphomas such as histiocytic lymphoma, anhydric hereditary ectodermal dysplasia, congenital alveolar dysplasia, and epithelial dysplasia of the cervix, fibrous dysplasia of bone, and mammary dysplasia, hyperplasias, e.g. endometrial, adrenal, breast, prostate or thyroid hyperplasias or pseudoepitheliomatous hyperplasia of

```
CC the skin.
XX
SQ Sequence 711 BP; 137 A; 214 C; 194 G; 159 T; 7 other;

  Query Match          90.6%; Score 15.4; DB 20; Length 711;
  Best Local Similarity 94.1%; Pred. No. 3.2e+02;
  Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 GGGAGCCCCCAGCAATGC 17
DB 478 GGGAGCCCCCAGCAAGC 494

RESULT 14
AA194530
ID AA194530 standard; cDNA; 785 BP.
XX
AC AA194530;
XX
DT 13-NOV-2001 (first entry)
XX
DE Human neuroblastoma expressed polynucleotide SEQ ID NO 605.
XX
KW Human; neuroblastoma; malignancy; cancer; tumour marker; N-myc; TrkA; ss.
XX
OS Homo sapiens.
XX
PN WO200166719-A1.
XX
PD 13-SEP-2001.
XX
PF 02-MAR-2001; 2001WO-JP01629.
XX
PR 07-MAR-2000; 2000JP-0159195.
XX
PA (CHIB-) CHIBA PREFECTURE.
PA (HISM) HISAMITSU PHARM CO LTD.
XX
PI Nakagawara A;
XX
WPI; 2001-565584/63.
XX
Nucleic acids originating in gene expressed in human neuroblastoma,
PT useful as probe or primer in diagnosing prognosis of human
PT neuroblastoma, malignancy and susceptibility indicator or tumour marker
PT for anti-cancer agents -
XX
Claim 1; Page 484-485; 2979pp; Japanese.
XX
The invention relates to novel genes (AA193926-AA197963) expressed in
CC human neuroblastoma. The nucleic acids are applicable as a probe or
CC primer in diagnosing the prognosis of human neuroblastoma, malignancy and
CC susceptibility indicators or tumour markers for anti-cancer agents. The
CC gene information for diagnosing prognosis is related to factors similar
CC to that for N-myc and TrkA genes.
XX
SQ Sequence 785 BP; 179 A; 261 C; 189 G; 135 T; 21 other;

  Query Match          90.6%; Score 15.4; DB 22; Length 785;
  Best Local Similarity 94.1%; Pred. No. 3.2e+02;
  Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 GGGAGCCCCCAGCAATGC 17
DB 359 GGGAGCCCCCAGCAATGC 375

RESULT 15
AA187832
ID AA187832 standard; cDNA; 805 BP.
XX
AC AA187832;
XX
```

```
DT 14-MAY-1998 (first entry)
XX
DE Human collagen 1A2 cDNA sequence containing the A polymorphism 907.
XX
KW Human; collagen 1A2; A polymorphism 907;
KW endogenous gene suppression; ss.
XX
OS Homo sapiens.
XX
PN WO9737014-A1.
XX
PD 09-OCT-1997.
XX
PF 02-APR-1997; 97WO-GB00929.
XX
PR 02-APR-1996; 96GB-0006961.
XX
PA (QUEEN-) QUEEN ELIZABETH COLLEGE DUBLIN.
XX
PI Farrar GJ, Humphries P, Kenna PF;
XX
WPI; 1997-503100/46.
XX
Endogenous gene suppression and replacement - useful in genetic
PT disease therapy
PT
PS Example 4; Page 82; 90pp; English.
XX
CC The present sequence was used in the development of a novel
CC strategy for suppressing at least part of the coding region of an
CC endogenous gene (tg), and replacing the suppressed gene sequence
CC with a nucleic acid sequence which differs from the EG. The
CC strategy can be useful when the gene, which is naturally present in
CC the genome of a patient, contributes to a disease state. Generally,
CC one allele of the gene will be mutated, i.e. it will possess
CC alterations in its nucleotide sequence that affect the function or
CC level of the gene product, e.g. the alteration may result in an
CC altered protein product from the wild type gene or control of
CC transcription and processing. Inheritance or somatic acquisition of
CC such a mutation can give rise, or predispose an individual to a
CC disease phenotype. However the gene can also be of wild type
CC phenotype, but contribute to a disease state in another way, so
CC that suppression would alleviate or improve the disease state or
CC improve the effectiveness of an administered therapeutic compound.
XX
SQ Sequence 805 BP; 179 A; 233 C; 162 G; 163 T; 68 other;

  Query Match          90.6%; Score 15.4; DB 18; Length 805;
  Best Local Similarity 94.1%; Pred. No. 3.2e+02;
  Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 GGGAGCCCCCAGCAATGC 17
DB 49 GGGAGCCCCCAGCAAGC 65

Search completed: January 21, 2003, 16:27:58
Job time : 144.611 secs
```

GenCore version 5.1.3  
Copyright (c) 1993 - 2003 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run On: January 21, 2003, 14:52:32 ; Search time 1340.03 Seconds  
(without alignments)  
229.633 Million cell updates/sec

Title: US-09-853-688-36  
Perfect score: 19  
Sequence: 1 ttaggaagtcgtgggtgc 19

Scoring table: IDENTITY\_NUC  
Gapop 10.0 , Gapext 1.0

Searched: 16154066 seqs, 8097743376 residues

Total number of hits satisfying chosen parameters: 32308132

Minimum DB seq length: 0  
Maximum DB seq length: 2000000000  
Post-processing: Minimum Match 0%  
Maximum Match 100%  
Listing first 45 summaries

Database :  
EST:  
1: em\_estba:\*  
2: em\_esthm:\*  
3: em\_estin:\*  
4: em\_estmu:\*  
5: em\_estov:\*  
6: em\_estpl:\*  
7: em\_estro:\*  
8: em\_hic:\*  
9: gb\_est1:\*  
10: gb\_est2:\*  
11: gb\_hic:\*  
12: gb\_est3:\*  
13: gb\_est4:\*  
14: gb\_est5:\*  
15: em\_estfun:\*  
16: em\_estom:\*  
17: gb\_gss:\*  
18: em\_gss\_hum:\*  
19: em\_gss\_inv:\*  
20: em\_gss\_pln:\*  
21: em\_gss\_vrt:\*  
22: em\_gss\_fun:\*  
23: em\_gss\_mam:\*  
24: em\_gss\_mus:\*  
25: em\_gss\_other:\*  
26: em\_gss\_pro:\*  
27: em\_gss\_rod:\*

pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

# SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
c 1	19	100.0	629	17	AQ047255
c 2	17	89.5	478	17	AZ165947
c 3	16.4	86.3	223	10	BB148197
c 4	16.4	86.3	245	10	BB365770
c 5	16.4	86.3	252	17	CNS048J2
c 6	16.4	86.3	302	10	BB041120

c 7	16.4	86.3	365	17	AQ062287
c 8	16.4	86.3	386	17	AZ739644
c 9	16.4	86.3	570	13	BI064819
c 10	16.4	86.3	660	17	AG074600
c 11	16.4	86.3	724	12	BE873650
c 12	16.4	86.3	974	17	CNS0444T
c 13	16	84.2	299	14	BQ256354
c 14	16	84.2	530	17	BH326060
c 15	16	84.2	908	17	AZ206811
c 16	16	84.2	1146	14	BM805318
c 17	15.8	83.2	187	9	AV016693
c 18	15.8	83.2	216	9	AV249326
c 19	15.8	83.2	225	9	AV141285
c 20	15.8	83.2	232	9	AT447750
c 21	15.8	83.2	261	17	AZ704742
c 22	15.8	83.2	279	9	AI585575
c 23	15.8	83.2	279	9	AV211876
c 24	15.8	83.2	289	10	BB469541
c 25	15.8	83.2	298	10	BE142089
c 26	15.8	83.2	355	12	BE138757
c 27	15.8	83.2	397	9	AA796032
c 28	15.8	83.2	401	9	AA138247
c 29	15.8	83.2	442	9	AA153692
c 30	15.8	83.2	444	9	AI645531
c 31	15.8	83.2	448	17	BH752980
c 32	15.8	83.2	484	12	BE918602
c 33	15.8	83.2	496	9	AI647277
c 34	15.8	83.2	508	13	BI811221
c 35	15.8	83.2	521	9	AI721115
c 36	15.8	83.2	532	13	BM220909
c 37	15.8	83.2	536	14	BQ394880
c 38	15.8	83.2	542	17	AZ860432
c 39	15.8	83.2	543	17	AZ883006
c 40	15.8	83.2	554	9	AU148749
c 41	15.8	83.2	586	10	AV958045
c 42	15.8	83.2	593	14	BQ745854
c 43	15.8	83.2	594	14	BP016690
c 44	15.8	83.2	599	17	AZ358149
c 45	15.8	83.2	612	17	AZ886074

## ALIGNMENTS

RESULT 1  
LOCUS AQ047255/c 629 bp DNA linear GSS 14-APR-1999  
DEFINITION RPC111-32118.TK RPC1-11 Homo sapiens genomic clone RPC1-11-32118, DNA sequence.  
ACCESSION AQ047255  
VERSION AQ047255.1 GI:3316182  
KEYWORDS GSS.  
SOURCE human.  
ORGANISM Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.  
REFERENCE 1 (bases 1 to 629)  
AUTHORS Adams,M.D., Rounsley,S.D., Zhao,S., Field,C.E., Bass,S., Linher,K., Golden,K., Berry,K., Granger,D., Suh,E., Wible,C., de Jong,P. and Venter,J.C.  
TITLE Use of BAC End Sequences for Sequence-Ready Map Building (1998)  
JOURNAL Unpublished (1998)  
COMMENT Contact: Mark Adams  
Department of Eukaryotic Genomics  
The Institute for Genomic Research  
9712 Medical Center Dr., Rockville, MD 20850, USA  
Tel: 301 838 0200  
Fax: 301 838 0208  
Email: mdadams@igrr.org  
Clones are derived from the human BAC library RPC1-11. For BAC library availability, please contact Pieter de Jong (pieter@dejong.med.buffalo.edu). Clones may be purchased from BACPAC Resources (http://bacpac.med.buffalo.edu/ordering) or from

Research Genetics (info@resgen.com). BAC end search page:

http://www.tigr.org/tdb/humgen/bac\_end\_search/bac\_end\_search.html  
 Class: BAC ends.

## FEATURES

## source

## Location/Qualifiers

1. .629  
 /organism="Homo sapiens"  
 /db\_xref="taxon:9606"  
 /clone="RPCI-11-32118"  
 /clone\_lib="RPCI-11"  
 /sex="Male"  
 /cell\_type="lymphocytes"  
 /note="Vector: pBACE3.6; Site\_1: EcoRI; Site\_2: EcoRI;  
 RPCI11 Human Male BAC Library"

## BASE COUNT

177 a 149 c 129 g 174 t

## ORIGIN

Query Match 100.0%; Score 19; DB 17; Length 629;

Best Local Similarity 100.0%; Pred. NO. 1.3e+02; Indels 0; Gaps 0;  
 Matches 19; Conservative 0; Mismatches 0;

QY 1 TGTAGGAAGTCTGGGTGC 19

DB 477 TGTAGGAAGTCTGGGTGC 459

## RESULT 2

## LOCUS

SP\_0083\_A2\_G02\_17A Strongylocentrotus purpuratus, purple sea urchin  
 clone Plate=83 Col=4 Row=M, DNA sequence.

## ACCESSION

AZ165947

## VERSION

AZ165947.1 GI:8335714

## KEYWORDS

## SOURCE

## ORGANISM

Strongylocentrotus purpuratus.

Eukaryota; Metazoa; Echinodermata; Eleutherozoa; Echinozoa;

Echinoidea; Euechinoidea; Echinacea; Echinoida;

Strongylocentrotidae; Strongylocentrotus.

1 (bases 1 to 478)

Cameron, R.A., Mahairas, G., Rast, J.P., Martinez, P., Biondi, T.R.,

Swartzell, S., Wallace, J.C., Foustka, A.J., Livingston, B.T., Wray

, G.A., Ettensohn, C.A., Lehrach, H., Britten, R.J., Davidson, E.H. and

Hood, L.

A sea urchin genome project: Sequence scan, virtual map, and

additional resources

PROC. Natl. Acad. Sci. U.S.A. 97 (17), 9514-9518 (2000)

Contact: Cameron, R.A., Davidson, E.H., Hood, L.

Division of Biology 156-29

California Institute of Technology

Pasadena California 91125, USA

Tel: (626) 395-8421

Fax: (626) 793-3047

Email: acameron@caltech.edu

Plate: 83 row: M column: 4

Seq primer: 17

Class: BAC ends

High quality sequence stop: 478.

Location/Qualifiers

1. .478

/organism="Strongylocentrotus purpuratus"

/db\_xref="taxon:7668"

/clone="Plate=83 Col=4 Row=M"

/clone\_lib="Strongylocentrotus purpuratus, purple sea

urchin, sperm genomic BAC library"

/note="Organ: sperm; Vector: BACE3.6; BAC Clones in E-Coli

DH10B"

150 a 93 c 88 g 140 t 7 others

Query Match 89.5%; Score 17; DB 17; Length 478;

Best Local Similarity 100.0%; Pred. No. 1.1e+03;

Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 TAGGAAGTCTGGGTGC 19

DB 53 TAGGAAGTCTGGGTGC 69

## RESULT 3

## LOCUS

BB148197 RIKEN full-length enriched, adult female vagina Mus  
 musculus cDNA clone 930106105 3', mRNA sequence.

## ACCESSION

## VERSION

## KEYWORDS

## SOURCE

## ORGANISM

## REFERENCE

## AUTHORS

## TITLE

## JOURNAL

## COMMENT

## CONTACT

## UNPUBLISHED

## LABORATORY

## SCIENCES CENTER

## THE INSTITUTE

## OF PHYSICAL

## AND CHEMICAL

## RESEARCH (RIKEN)

## 1-7-22

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## Tsurumi-ku,

## Yokohama,

## Kanagawa

## 230-0045, Japan

## Tel: 81-45-503-9222

## Fax: 81-45-503-9216

## Email: genome.res@gsic.riken.go.jp,

## URL: http://genome.gsc.riken.go.jp/

## Carninci, P., Nishiyama, Y., Westover, A., Itoh, M., Nagaoka, S., Sasaki

## N., Okazaki, Y., Muramatsu, M. and Hayashizaki, Y.

## Thermotabilization and thermoactivation of thermolabile enzymes by

## trehalose and its application for the synthesis of full length

## cDNA. Proc. Natl. Acad. Sci. U.S.A. 95 (2), 520-524 (1998)

## Itoh, M., Katsunai, T., Akiyama, J., Shibata, K., Izawa, M., Kawai, J.,

## Tomaru, Y., Carninci, P., Shibata, Y., Ozawa, Y., Muramatsu, M., Okazaki

## , Y. and Hayashizaki, Y.

## Automated filtration-based high-throughput plasmid preparation

## system. Genome Res. 9 (5), 463-470 (1999)

## Carninci, P. and Hayashizaki, Y.

## High-efficiency full-length cDNA cloning. Methods Enzymol. 303,

## 19-44 (1999)

## Please visit our web site (http://genome.rtc.riken.go.jp) for

## further details.

## Location/Qualifiers

## 1. .223

## /organism="Mus musculus"

## /db\_xref="taxon:10090"

## /clone="930106105"

## /clone\_lib="RIKEN full-length enriched, adult female

## vagina"

## /sex="female"

## /tissue\_type="vagina"

## /dev\_stage="adult"

## /lab\_host="DH10B"

## /note="Site 1: SalI; Site 2: BamHI; cDNA library was

## prepared and sequenced in Mouse Genome Encyclopedia

## Project of Genome Exploration Research Group in Riken

Genomic Sciences Center and Genome Science Laboratory in RIKEN. Division of Experimental Animal Research in Riken contributed to prepare mouse tissues. 1st strand cDNA was primed with a primer [5'.

GAGAGAGAGAGATCCAGAGCTCTTTTTTTTTTTTNN 3'], cDNA was prepared by using trehalose thermo-activated reverse transcriptase and subsequently enriched for full-length by cap-trapper. cDNA went through one round of normalization to kot -10.0 and subtraction to Rot -185.0. Second strand cDNA was prepared with the primer adapter of sequence [5' GAGAGAGAGATTCGAGTTAAATTAATCCCCCCCCCC 3']. cDNA was cleaved with XhoI and BamHI. Vector: a modified pBluescript KS(+) after bulk excision from Lambda FUC I."

BASE COUNT 50 a 72 c 35 g 66 t  
 ORIGIN  
 Query Match 86.3%; Score 16.4; DB 10; Length 223;  
 Best Local Similarity 94.4%; Pred. NO. 1.8e+03;  
 Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 2 GTAGGAAGTCTGGGGTGC 19  
 ||||| ||||| |||||

DB 77 GTAGGAAGTCTGAGGTGC 60

RESULT 4  
 BB365770/c  
 LOCUS  
 DEFINITION BB365770 RIKEN full-length enriched, 16 days embryo head Mus  
 musculus cDNA clone C130030N17 3', mRNA sequence.

ACCESSION BB365770.1 GI:9077598  
 VERSION  
 KEYWORDS EST.  
 SOURCE house mouse.  
 ORGANISM Mus musculus

REFERENCE  
 AUTHORS Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

1 (bases 1 to 245)  
 Konno,H., Alzawa,K., Akahira,S., Akiyama,J., Arakawa,T., Carninci,P., Endo,T., Fukuda,S., Fukunishi,Y., Hara,A., Hayatsu,N., Hirozane,T., Hori,F., Ishii,Y., Ishikawa,J., Ishikawa,T., Itoh,M., Izawa,M., Kadota,K., Kagawa,I., Kai,C., Kawai,J., Kikuchi,N., Kiyosawa,H., Kojima,Y., Kondo,S., Koya,S., Kurihara,C., Kusakabe,M., Matsuyama,T., Miki,R., Mizuno,Y., Nakamura,M., Oda,H., Okazaki,Y., Ono,T., Owa,C., Saito,H., Sakai,C., Sato,K., Shibata,K., Shibata,Y., Shigemoto,Y., Shinagawa,A., Shiraki,T., Sogabe,Y., Sugahara,Y., Suzuki,H., Suzuki,H., Tagawa,A., Takahashi,F., Tomimaga,N., Toya,T., Tsunoda,Y., Watanabe,S., Watanabe,S., Yamamura,T., Yamanaka,I., Yano,R., Yasunishi,A., Yokota,T., Yoshida,K., Yoshiki,A., Yoshino,M., Muramatsu,M. and Hayashizaki,Y.  
 RIKEN Mouse ESTs (Konno.H., et al.)  
 Unpublished (2000)

TITLE  
 JOURNAL  
 COMMENT Contact: Yoshihide Hayashizaki  
 Laboratory for Genome Exploration Research Group, RIKEN Genomic Sciences Center(GSC) Yokohama Institute  
 The Institute of Physical and Chemical Research (RIKEN)  
 1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan  
 Tel: 81-45-503-9222  
 Fax: 81-45-503-9216  
 Email: genome-res@sc.riken.go.jp,  
 URL:http://genome.gsc.riken.go.jp/

Carninci,P., Nishiyama,Y., Westover,A., Itoh,M., Nagaoka,S., Sasaki,N., Okazaki,Y., Muramatsu,M. and Hayashizaki,Y.  
 Thermoactivation and thermoactivation of thermolabile enzymes by trehalose and its application for the synthesis of full length cDNA. Proc. Natl. Acad. Sci. U.S.A. 95 (2), 520-524 (1998)  
 Itoh,M., Kitsunai,T., Akiyama,J., Shibata,K., Izawa,M., Kawai,J., Tomaru,Y., Carninci,P., Shibata,Y., Ozawa,Y., Muramatsu,M., Okazaki,Y. and Hayashizaki,Y.  
 Automated filtration-based high-throughput plasmid preparation system. Genome Res. 9 (5), 463-470 (1999)  
 Carninci,P. and Hayashizaki,Y.

High-efficiency full-length cDNA cloning. Methods Enzymol. 303, 19-44 (1999)  
 Please visit our web site (<http://genome.rtc.riken.go.jp>) for further details.

FEATURES  
 source

Location/Qualifiers  
 1..245  
 /organism="Mus musculus"  
 /strain="C57BL/6J"  
 /db\_xref="taxon:10090"  
 /clone="C130030N17"  
 /clone\_lib="RIKEN full-length enriched, 16 days embryo head"  
 /sex="mixed"  
 /tissue\_type="head"  
 /dev\_stage="16 days embryo"  
 /lab\_host="DH10B"  
 /note="Site\_1: SalI; Site\_2: BamHI; cDNA library was prepared and sequenced in Mouse Genome Encyclopedia Project of Genome Exploration Research Group in Riken Genomic Sciences Center and Genome Science Laboratory in RIKEN. Division of Experimental Animal Research in Riken contributed to prepare mouse tissues. 1st strand cDNA was primed with a primer [5' GAGAGAGAGATCCAGAGCTCTTTTTTTTTTTTNN 3'], cDNA was prepared by using trehalose thermo-activated reverse transcriptase and subsequently enriched for full-length by cap-trapper. Second strand cDNA was prepared with the primer adapter of sequence [5' GAGAGAGATTCGAGTTAAATTAATTAATCCCCCCCCCC 3']. cDNA was cloned into the XhoI and BamHI sites. Vector: a modified pBluescript KS(+) after bulk excision from Lambda FUC I"

BASE COUNT 76 a 80 c 43 g 46 t  
 ORIGIN

Query Match 86.3%; Score 16.4; DB 10; Length 245;  
 Best Local Similarity 94.4%; Pred. No. 1.8e+03;  
 Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 2 GTAGGAAGTCTGGGGTGC 19  
 ||||| ||||| |||||

DB 50 GTAGGAATCTGGGGTGC 33

RESULT 5  
 CNS04BJ2/c  
 LOCUS

DEFINITION CNS04BJ2 252 bp DNA linear GSS 21-MAY-2000  
 Tetraodon nigroviridis genome survey sequence PUC-ori end of clone 097H01 of library G from Tetraodon nigroviridis, genomic survey sequence.

ACCESSION AL283223  
 VERSION AL283223.1 GI:8021581  
 KEYWORDS GSS: genome survey sequence.  
 SOURCE Tetraodon nigroviridis.

ORGANISM Tetraodon nigroviridis

REFERENCE  
 AUTHORS Roest-Crollius,H., Jaillon,O., Dasilva,C., Bouneau,L., Fisher,C., Bernot,A., Fizames,C., Wincker,P., Brottier,P., Quetier,F., Saurin,W. and Weissenbach,J.  
 Human gene number estimate provided by genome wide analysis using Tetraodon nigroviridis DNA sequence

TITLE 1 (bases 1 to 252)

JOURNAL 2 (bases 1 to 252)  
 REFERENCE Roest-Crollius,H., Jaillon,O., Dasilva,C., Fizames,C., Fisher,C., Bouneau,L., Billault,A., Quetier,F., Saurin,W., Bernot,A. and Weissenbach,J.  
 Characterization and repeat analysis of the compact genome of the freshwater pufferfish Tetraodon nigroviridis

JOURNAL Unpublished

TITLE Unpublished

JOURNAL Unpublished





Email: mdadamst@tigr.org  
Clones are available from Research Genetics (info@resgen.com). BAC  
end search page:  
[http://www.tigr.org/tdb/humgen/bac\\_end\\_search.html](http://www.tigr.org/tdb/humgen/bac_end_search.html).  
seq primer: M13-21

## FEATURES

```

BASE COUNT      97 a      88 q      94 t
ORIGIN
      86.3%   Score 16.4;   DB 17:   length 365;
Query Match     94.4%;
Best Local Similarity
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

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	FEATURES	SOURCE
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100.	... ..	...

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Location/Qualifiers
1. 386
/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="RPCI-24-71H2"
/clone_lib="RPCI-24"
/sex="Male"
/cell_type="Spleen/Brain"
/note="vector: P1RHHAC1; Site_1: BamHI; Site_2: BamHI;
RPCI-24 Mouse BAC Library produced by Pieter de Jong. The

```

Department of Eukaryotic Genomics  
The Institute for Genomic Research  
9712 Medical Center Dr., Rockville, MD 20850, USA  
Tel: 301 838 0200  
Fax: 301 838 0208  
Email: szhao@tigr.org  
Clones are derived from the mouse BAC library RPCI-24. For BAC library availability, please contact Pieter de Jong (pjejong@email.cho.org). Clones may be purchased from BACPAC Resources (<http://www.choi.org/bacpac/orderingframe.htm>). BAC end page: [http://www.tigr.org/tdb/bac\\_ends/mouse/bac\\_end\\_intro.html](http://www.tigr.org/tdb/bac_ends/mouse/bac_end_intro.html)  
Plate: 71 row: H column: 22



```

UY 1 TGTAGGAAGTCTGGGTG 18
Db 457 TGTAGGAAGTCTGAGGTG 340

RESULT 13
LOCUS BH256354 299 bp mRNA linear EST 06-MAY-2002
DEFINITION NTSC_j012g04.y1 NCI_CGAP_BCC2 Mus musculus cDNA clone IMAGE:4113607
5' mRNA sequence.
ACCESSION BH256354
VERSION RQ256354.1 GI:20457107
KEYWORDS EST.
SOURCE house mouse.
ORGANISM Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Mus.
1 (bases 1 to 299)
NCI-CCAP http://www.ncbi.nlm.nih.gov/ncicgap.
National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index
Unpublished (1997)
JOURNAL Contact: Robert Strausberg, Ph.D.
COMMENT Email: cgaops@mail.nih.gov
cDNA Library Preparation: David B. Krizman, Ph.D.
cDNA Library Arrayed by: The I.M.A.G.E. Consortium/LLNL
DNA Sequencing by: National Institutes of Health Intramural
Sequencing Center (NISC)
Clone distribution: NCI-CCAP clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
info@image.llnl.gov
MGI:1538759
Plate: LHAM9369 row: N column: 8
Seq primer: M13RPI reverse primer (ABI).
FEATURES
Location/Qualifiers
1..299
/organism="Mus musculus"
/db_xref="taxon:10090"
/clone="IMAGE:4113607"
/clone_lib="NCI_CGAP_BCC2"
/tissue_type="flow-sorted, common granulocyte-macrophage
progenitors"
/lab_host="DH10B"
/notes="Organ: blood; Vector: pAMP1; mRNA made from
granulocyte-macrophage progenitors, cDNA made by oligo-dT
priming. Directionally cloned into UDG sites.
Size-selected on agarose gel, average insert size 300 bp.
Primary library. cDNA Library Preparation: David B.
Krizman, Ph.D."
BASE COUNT 77 a 71 c 96 g 55 t
ORIGIN
Query Match 84.2%; Score 16; DB 14; Length 299;
Best Local Similarity 100.0%; Pred. No. 2.8e+03;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 4 AGGAAGTCTGGGTGC 19
|||||
Db 120 AGGAAGTCTGGGTGC 135

RESULT 14
LOCUS BH326060 530 bp DNA linear GSS 03-DEC-2001
DEFINITION CH230-93G3.TJ CHORI-230 Segment 1 Rattus norvegicus genomic clone
CH230-93G3, DNA sequence.
ACCESSION BH326060
VERSION BH326060.1 GI:17256774
KEYWORDS GSS.
SOURCE Norway rat.
ORGANISM Rattus norvegicus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;

```

```

Rattus.
1 (bases 1 to 530)
Zhao, S., Shetty, J., Shatsman, S., Tsegaye, G., Geer, K., Shvartsbeyn
, A., Gebregorgis, E., Overton, L., Russell, D., Chen, D., Riggs, F., de
Jong, P., and Fraser, C. M.
Rat HAC End Sequences from Library CHORI-230 EcoRI segment
Unpublished (1999)
Other_GSSs: CH230-93G3.TV
Contact: Shaying Zhao
Department of Eukaryotic Genomics
The Institute for Genomic Research
9712 Medical Center Dr., Rockville, MD 20850, USA
Tel: 301 838 0200
Fax: 301 838 0208
Email: szhao@tigr.org
Clones are derived from the rat BAC library CHORI-230
(http://www.chori.org/bacpac/rat230.htm). For BAC library
availability, please contact Pieter de Jong (pdejong@mail.cho.org).
Clones may be purchased from BACPAC Resources
(http://www.chori.org/bacpac/or ering_information.htm). BAC end
page: http://www.tigr.org/tldb/bac_ends/rat/bac_end_intro.html
Plate: 93 row: G column: 3
Seq primer: SP6
Class: BAC ends.
Location/Qualifiers
1..530
/organism="Rattus norvegicus"
/strain="BN/SSNHsd/MCW"
/db_xref="taxon:10116"
/clone="CH230-93G3"
/clone_lib="CHORI-230 Segment 1"
/sex="Female"
/cell_type="Brain"
/notes="Vector: pTARBAC2.1; Site_1: EcoRI; Site_2: EcoRI;
CHORI-230 Rat (BN/SSNHsd/MCW) BAC library produced by
Pieter de Jong"
BASE COUNT 123 a 130 c 117 g 160 t
ORIGIN
Query Match 84.2%; Score 16; DB 17; Length 530;
Best Local Similarity 100.0%; Pred. No. 3.1e+03;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 4 AGGAAGTCTGGGTGC 19
|||||
Db 74 AGGAAGTCTGGGTGC 89

RESULT 15
LOCUS AZ206811/c 908 bp DNA linear GSS 31-AUG-2000
DEFINITION SP_0109_AL_H02_T7A Strongylocentrotus purpuratus, purple sea urchin
, sperm genomic BAC library Strongylocentrotus purpuratus genomic
clone Plate-109 Col-3 Row=O, DNA sequence.
ACCESSION AZ206811
VERSION AZ206811.1 GI:8417759
KEYWORDS GSS.
SOURCE Strongylocentrotus purpuratus.
ORGANISM Strongylocentrotus purpuratus
Strongylocentrotus purpuratus
Eukaryota; Metazoa; Echinodermata; Eleutherozoa; Echinozoa;
Echinoidea; Euechinoidea; Echinacea; Echinoida;
Strongylocentrotidae; Strongylocentrotus.
1 (bases 1 to 908)
Cameron, R.A., Mahairas, G., Rast, J.P., Martinez, P., Blondi, T.R.,
Swartzell, S., Wallace, J.C., Pousta, A.J., Livingston, B.T., Wray
, G.A., Ettensohn, C.A., Lehrach, H., Britten, R.J., Davidson, E.H. and
Hood, L.
A sea urchin genome project: Sequence scan, virtual map, and
additional resources
Proc. Natl. Acad. Sci. U.S.A. 97 (17), 9514-9518 (2000)
20402566
JOURNAL MEDLINE
COMMENT Contact: Cameron, R.A., Davidson, E.H., Hood, L
Division of Biology 156-29

```

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Pasadena California 91125, USA  
Tel: (626) 395-8421  
Fax: (626) 793-3047  
Email: acameron@caltech.edu  
Plate: 109 row: 0 column: 3  
Seq primer: 47  
Class: BAC ends  
High quality sequence stop: 908.

FEATURES  
Source  
1. .908  
location/Qualifiers  
/organism="Strongylocentrotus purpuratus"  
/db\_xref="taxon:7668"  
/clone="plate-109 Col-3 Row=0"  
/clone\_lib="Strongylocentrotus purpuratus, purple sea  
urchin, sperm genomic BAC library"  
/note="Organ: sperm; Vector: BACe3.6; BAC Clones in E-Coli  
DH10b"

BASE COUNT 226 a 194 c 169 g 319 t  
ORIGIN

Query Match 84.2%; Score 16; db 17; Length 908;  
Best Local Similarity 100.0%; Pred. No. 3.3e+03;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 4 AGGAAGTCTGGGTGC 19  
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db 344 AGGAAGTCTGGGTGC 319

Search completed: January 21, 2003, 15:38:08  
Job time : 1345.03 secs



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; FILE REFERENCE: WCM78
; CURRENT APPLICATION NUMBER: US/09/853,688
; CURRENT FILING DATE: 2001-05-14
; NUMBER OF SEQ ID NOS: 66
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 5
; LENGTH: 3700
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-853-688-5

Query Match      100.0%; Score 19; DB 10; Length 3700;
Best Local Similarity 100.0%; Pred. No. 1.4; 0; Indels 0; Gaps 0;
Matches 19; Conservative 0; Mismatches 0;

QY 1 TGTAGGAAGTCTGGGTCG 19
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Db 3316 TGTAGGAAGTCTGGGTCG 3298

RESULT 3
US-09-728-446-1120/c
; Sequence 1120, Application US/09728446
; Patent No. US20020081668A1
; GENERAL INFORMATION:
; APPLICANT: Friedrich, Glenn
; APPLICANT: Zambrowicz, Brian
; APPLICANT: Sands, Arthur T.
; TITLE OF INVENTION: No. US20020081668A1 Murine Polynucleotide Sequences
; FILE REFERENCE: LEX-0101-USA
; CURRENT APPLICATION NUMBER: US/09/728,446
; PRIOR FILING DATE: 2000-11-30
; PRIOR APPLICATION NUMBER: US 60/168,270
; PRIOR FILING DATE: 1999-12-01
; NUMBER OF SEQ ID NOS: 1461
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 1120
; LENGTH: 232
; TYPE: DNA
; ORGANISM: Mus musculus
; FEATURE:
; NAME/KEY: misc.feature
; LOCATION: (1)...(232)
; OTHER INFORMATION: n = A,T,C or G
US-09-728-446-1120

Query Match      83.2%; Score 15.8; DB 10; Length 232;
Best Local Similarity 89.5%; Pred. No. 4.4;
Matches 17; Conservative 0; Mismatches 2; Indels 2; Gaps 0;

QY 1 TGTAGGAAGTCTGGGTCG 19
   |||
Db 144 TGAAGGAAGTCTGGGTCG 126

RESULT 4
US-09-867-701-388/c
; Sequence 388, Application US/09867701
; Patent No. US2002013237A1
; GENERAL INFORMATION:
; APPLICANT: Aglate, Paul A.
; APPLICANT: Jones, Robert
; APPLICANT: Harlocker, Susan L.
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR THE THERAPY
; FILE REFERENCE: 210121.497
; CURRENT APPLICATION NUMBER: US/09/867,701
; CURRENT FILING DATE: 2001-05-29
; NUMBER OF SEQ ID NOS: 10912
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 388
; LENGTH: 170
; TYPE: DNA
; ORGANISM: Homo sapiens

; FILE REFERENCE: WCM78
; CURRENT APPLICATION NUMBER: US/09/853,688
; CURRENT FILING DATE: 2001-05-14
; NUMBER OF SEQ ID NOS: 66
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 5
; LENGTH: 3700
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-853-688-5

Query Match      81.1%; Score 15.4; DB 10; Length 170;
Best Local Similarity 94.1%; Pred. No. 67;
Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 2 GTAGGAAGTCTGGGTCG 18
   |||
Db 115 GTGGGAAGTCTGGGTCG 99

RESULT 5
US-09-864-761-11292/c
; Sequence 11292, Application US/09864761
; Patent No. US20020048763A1
; GENERAL INFORMATION:
; APPLICANT: Penn, Sharron G.
; APPLICANT: Hanzel, David K.
; APPLICANT: Chen, Wensheng
; TITLE OF INVENTION: HUMAN GENOME-DERIVED SINGLE EXON NUCLEIC ACID PROBES USEFUL FOR
; FILE REFERENCE: Aeonica-X-1
; CURRENT APPLICATION NUMBER: US/09/864,761
; CURRENT FILING DATE: 2001-05-23
; PRIOR FILING DATE: 2000-02-04
; PRIOR APPLICATION NUMBER: US 60/180,312
; PRIOR FILING DATE: 2000-05-26
; PRIOR APPLICATION NUMBER: US 09/632,366
; PRIOR FILING DATE: 2000-08-03
; PRIOR APPLICATION NUMBER: GB 24263.6
; PRIOR FILING DATE: 2000-10-04
; PRIOR APPLICATION NUMBER: US 60/236,359
; PRIOR FILING DATE: 2000-09-27
; PRIOR APPLICATION NUMBER: PCT/US01/006666
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00667
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00664
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00669
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00665
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00668
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00663
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00662
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00661
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00670
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: US 60/234,687
; PRIOR FILING DATE: 2000-09-21
; PRIOR APPLICATION NUMBER: US 09/608,408
; PRIOR FILING DATE: 2000-06-30
; PRIOR APPLICATION NUMBER: US 09/774,203
; PRIOR FILING DATE: 2001-01-29
; NUMBER OF SEQ ID NOS: 49117
; SOFTWARE: Annomax Sequence Listing Engine vers. 1.1
; SEQ ID NO 11292
; LENGTH: 456
; TYPE: DNA
; ORGANISM: Homo sapiens
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; FEATURE:
; OTHER INFORMATION: MAP TO AC005342.1
; OTHER INFORMATION: EXPRESSED IN LUNG, SIGNAL ~ 0.84
; OTHER INFORMATION: EXPRESSED IN BRAIN, SIGNAL ~ 1.1
; OTHER INFORMATION: EXPRESSED IN BONE MARROW, SIGNAL ~ 1.2
; OTHER INFORMATION: EXPRESSED IN PLACENTA, SIGNAL ~ 0.73
; OTHER INFORMATION: EXPRESSED IN FETAL LIVER, SIGNAL ~ 0.94
; OTHER INFORMATION: EXPRESSED IN ADULT LIVER, SIGNAL ~ 0.76
US-09-864-761-11292

Query Match      81.1%; Score 15.4; DB 10; Length 456;
Best Local Similarity 94.1%; Pred. No. 73;
Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TGTAGGAAGTCTGGGCT 17
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Db 125 TGTAGGAAGTCTGGGCT 109

RESULT 6
US-09-917-800A-410
; Sequence 410, Application US/09917800A
; Patent No. US20020119462A1
; GENERAL INFORMATION:
; APPLICANT: Mendrick, Donna
; APPLICANT: Porter, Mark
; APPLICANT: Johnson, Kory
; APPLICANT: Castle, Arthur
; APPLICANT: Elashoff, Michael
; TITLE OF INVENTION: Molecular Toxicology Modeling
; FILE REFERENCE: 44921-5038-US
; CURRENT APPLICATION NUMBER: US/09/917,800A
; CURRENT FILING DATE: 2001-07-31
; PRIOR APPLICATION NUMBER: US 60/222,040
; PRIOR FILING DATE: 2000-07-31
; PRIOR APPLICATION NUMBER: US 60/222,880
; PRIOR FILING DATE: 2000-11-02
; PRIOR APPLICATION NUMBER: US 60/290,029
; PRIOR FILING DATE: 2001-05-11
; PRIOR APPLICATION NUMBER: US 60/290,645
; PRIOR FILING DATE: 2001-05-15
; PRIOR APPLICATION NUMBER: US 60/292,336
; PRIOR FILING DATE: 2001-05-22
; PRIOR APPLICATION NUMBER: US 60/295,798
; PRIOR FILING DATE: 2001-06-06
; PRIOR APPLICATION NUMBER: US 60/297,457
; PRIOR FILING DATE: 2001-06-13
; PRIOR APPLICATION NUMBER: US 60/298,884
; PRIOR FILING DATE: 2001-06-19
; PRIOR APPLICATION NUMBER: US 60/303,459
; NUMBER OF SEQ ID NOS: 1740
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 410
; LENGTH: 392
; TYPE: DNA
; ORGANISM: Rattus norvegicus
; FEATURE:
; OTHER INFORMATION: Genbank Accession No. US20020119462A1 U10357
US-09-917-800A-410

Query Match      78.9%; Score 15; DB 10; Length 392;
Best Local Similarity 100.0%; Pred. No. 1.1e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 TAGGAAGTCTGGGCT 17
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Db 2036 TAGGAAGTCTGGGCT 2022

RESULT 8
US-09-764-868-1347/c
; Sequence 1347, Application US/09764868
; Patent No. US20020168711A1
; GENERAL INFORMATION:
; APPLICANT: Rosen et al.
; TITLE OF INVENTION: Nucleic Acids, Proteins, and Antibodies
; FILE REFERENCE: PT232
; CURRENT APPLICATION NUMBER: US/09/764,868
; CURRENT FILING DATE: 2001-01-17
; Prior application data removed - refer to PALM or file wrapper
; NUMBER OF SEQ ID NOS: 1510
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 1347
; LENGTH: 24757
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-764-868-1347

Query Match      78.9%; Score 15; DB 9; Length 24757;
Best Local Similarity 100.0%; Pred. No. 1.6e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 TAGGAAGTCTGGGCT 17
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Db 190 TAGGAAGTCTGGGCT 204

RESULT 7
US-09-917-800A-1449/c
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; Sequence 1449, Application US/09917800A
; Patent No. US20020119462A1
; GENERAL INFORMATION:
; APPLICANT: Mendrick, Donna
; APPLICANT: Porter, Mark
; APPLICANT: Johnson, Kory
; APPLICANT: Castle, Arthur
; APPLICANT: Elashoff, Michael
; APPLICANT: Gene Logic, Inc.
; TITLE OF INVENTION: Molecular Toxicology Modeling
; FILE REFERENCE: 44921-5038-US
; CURRENT APPLICATION NUMBER: US/09/917,800A
; CURRENT FILING DATE: 2001-07-31
; PRIOR APPLICATION NUMBER: US 60/222,040
; PRIOR FILING DATE: 2000-07-31
; PRIOR APPLICATION NUMBER: US 60/222,880
; PRIOR FILING DATE: 2000-11-02
; PRIOR APPLICATION NUMBER: US 60/290,029
; PRIOR FILING DATE: 2001-05-11
; PRIOR APPLICATION NUMBER: US 60/290,645
; PRIOR FILING DATE: 2001-05-15
; PRIOR APPLICATION NUMBER: US 60/292,336
; PRIOR FILING DATE: 2001-05-22
; PRIOR APPLICATION NUMBER: US 60/295,798
; PRIOR FILING DATE: 2001-06-06
; PRIOR APPLICATION NUMBER: US 60/297,457
; PRIOR FILING DATE: 2001-06-13
; PRIOR APPLICATION NUMBER: US 60/298,884
; PRIOR FILING DATE: 2001-06-19
; PRIOR APPLICATION NUMBER: US 60/303,459
; NUMBER OF SEQ ID NOS: 1740
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 1449
; LENGTH: 2207
; TYPE: DNA
; ORGANISM: Rattus norvegicus
; FEATURE:
; OTHER INFORMATION: Genbank Accession No. US20020119462A1 U10357
US-09-917-800A-1449

Query Match      78.9%; Score 15; DB 10; Length 2207;
Best Local Similarity 100.0%; Pred. No. 1.3e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 TAGGAAGTCTGGGCT 17
   ||||| ||||| |||||
Db 2036 TAGGAAGTCTGGGCT 2022

RESULT 8
US-09-764-868-1347/c
; Sequence 1347, Application US/09764868
; Patent No. US20020168711A1
; GENERAL INFORMATION:
; APPLICANT: Rosen et al.
; TITLE OF INVENTION: Nucleic Acids, Proteins, and Antibodies
; FILE REFERENCE: PT232
; CURRENT APPLICATION NUMBER: US/09/764,868
; CURRENT FILING DATE: 2001-01-17
; Prior application data removed - refer to PALM or file wrapper
; NUMBER OF SEQ ID NOS: 1510
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 1347
; LENGTH: 24757
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-764-868-1347

Query Match      78.9%; Score 15; DB 9; Length 24757;
Best Local Similarity 100.0%; Pred. No. 1.6e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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QY 5 GCAAGTCTGGGGTGC 19  
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DB 10501 GCAAGTCTGGGGTGC 10487

## RESULT 9

US-09-983-965-2137/c  
; Sequence 2137, Application US/09983965  
; Patent No. US20020137160A1  
; GENERAL INFORMATION:  
; APPLICANT: Warren, Wesley C.  
; APPLICANT: Tao, Nengbing  
; APPLICANT: Ryatt, John C.  
; APPLICANT: Mathalagan, Nadappan  
; TITLE OF INVENTION: NUCLEIC ACID AND OTHER MOLECULES ASSOCIATED WITH LACTATION AND  
; FILE OF INVENTION: MUSCLE AND FAT DEPOSITION  
; FILE REFERENCE: 37-21(10297)C  
; CURRENT APPLICATION NUMBER: US/09/983,965  
; CURRENT FILING DATE: 2001-10-26  
; PRIOR APPLICATION NUMBER: US 09/465,231  
; PRIOR FILING DATE: 1999-12-15  
; PRIOR APPLICATION NUMBER: US 60/113,678  
; PRIOR FILING DATE: 1998-12-17  
; NUMBER OF SEQ ID NOS: 5912  
; SEQ ID NO 2137  
; LENGTH: 317  
; TYPE: DNA  
; ORGANISM: Bos taurus  
; FEATURE:  
; OTHER INFORMATION: Clone ID: 37-LIB3057-014-Q1-K1-R2  
US-09-983-965-2137

Query Match 77.9%; Score 14.8; DB 10; Length 317;  
Best Local Similarity 88.9%; Pred. No. 1.4e+02;  
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 TGTAGGAAGTCTGGGGTGC 18  
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DB 200 TGAAGGAAGTCTGGGGTGC 183

## RESULT 10

US-09-925-300-178/c  
; Sequence 178, Application US/09925300  
; Patent No. US20020151681A1  
; GENERAL INFORMATION:  
; APPLICANT: Craig Rosen,  
; APPLICANT: Steve Ruben,  
; TITLE OF INVENTION: Nucleic Acids, Proteins and Antibodies  
; FILE REFERENCE: PA101  
; CURRENT APPLICATION NUMBER: US/09/925,300  
; CURRENT FILING DATE: 2001-08-10  
; PRIOR APPLICATION NUMBER: PCT/US00/05988  
; PRIOR FILING DATE: 2000-03-08  
; PRIOR APPLICATION NUMBER: 60/124,270  
; PRIOR FILING DATE: 1999-03-12  
; NUMBER OF SEQ ID NOS: 1890  
; SOFTWARE: Patentin Ver. 2.0  
; SEQ ID NO 178  
; LENGTH: 393  
; TYPE: DNA  
; ORGANISM: Homo sapiens  
; FEATURE:  
; NAME/KEY: misc feature  
; LOCATION: (214)  
; OTHER INFORMATION: n equals a,t,g, or c  
US-09-925-300-178

Query Match 77.9%; Score 14.8; DB 10; Length 393;  
Best Local Similarity 84.2%; Pred. No. 1.4e+02;  
Matches 16; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 TGTAGGAAGTCTGGGGTGC 19

DB 224 TGTAGCAAGTNAGGGGTGC 206  
||||| ||||| |||||

## RESULT 11

US-09-864-761-16125/c  
; Sequence 16125, Application US/09864761  
; Patent No. US20020048763A1  
; GENERAL INFORMATION:  
; APPLICANT: Penn, Sharron G.  
; APPLICANT: Rank, David R.  
; APPLICANT: Hanzel, David K.  
; APPLICANT: Chen, Wensheng  
; TITLE OF INVENTION: HUMAN GENOME-DERIVED SINGLE EXON NUCLEIC ACID PROBES USEFUL FO  
; FILE REFERENCE: Acomica-X-1  
; CURRENT APPLICATION NUMBER: US/09/864,761  
; CURRENT FILING DATE: 2001-05-23  
; PRIOR APPLICATION NUMBER: US 60/180,312  
; PRIOR FILING DATE: 2000-02-04  
; PRIOR APPLICATION NUMBER: US 60/207,456  
; PRIOR FILING DATE: 2000-05-26  
; PRIOR APPLICATION NUMBER: US 09/632,366  
; PRIOR FILING DATE: 2000-08-03  
; PRIOR APPLICATION NUMBER: GB 24263.6  
; PRIOR FILING DATE: 2000-10-04  
; PRIOR APPLICATION NUMBER: US 60/236,359  
; PRIOR FILING DATE: 2000-09-27  
; PRIOR APPLICATION NUMBER: PCT/US01/00666  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/00667  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/00664  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/00669  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/00665  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/00668  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/00663  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/00662  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/00661  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/00670  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: US 60/234,687  
; PRIOR FILING DATE: 2000-09-21  
; PRIOR APPLICATION NUMBER: US 09/608,408  
; PRIOR FILING DATE: 2000-06-30  
; PRIOR APPLICATION NUMBER: US 09/774,203  
; PRIOR FILING DATE: 2001-01-29  
; NUMBER OF SEQ ID NOS: 49117  
; SOFTWARE: Annomax Sequence Listing Engine vers. 1.1  
; SEQ ID NO 16125  
; LENGTH: 558  
; TYPE: DNA  
; ORGANISM: Homo sapiens  
; FEATURE:  
; OTHER INFORMATION: MAP TO AC010486.5  
; OTHER INFORMATION: EXPRESSED IN PLACENTA, SIGNAL = 0.47  
; OTHER INFORMATION: EXPRESSED IN BONE MARROW, SIGNAL = 0.46  
US-09-864-761-16125

Query Match 77.9%; Score 14.8; DB 10; Length 558;  
Best Local Similarity 88.9%; Pred. No. 1.5e+02;  
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 GTAGGAAGTCTGGGGTGC 19  
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DB 362 GCAGGAAGTCAGGGGTGC 345



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: GENERAL INFORMATION:
: APPLICANT: Rosen et al.
: TITLE OF INVENTION: Nucleic Acids, Proteins, and Antibodies
: FILE REFERENCE: P1110
: CURRENT APPLICATION NUMBER: US/09/764,855
: CURRENT FILING DATE: 2001-01-17
: Prior application data removed - consult PALM or file wrapper
: NUMBER OF SEQ ID NOS: 334
: SOFTWARE: Patentin Ver. 2.0
: SEQ ID NO 202
: LENGTH: 3253
: TYPE: DNA
: ORGANISM: Homo sapiens
US-09-764-855-202

Query Match      77.9%; Score 14.8; DB 10; Length 3253;
Best Local Similarity 88.9%; Pred. No. 1.7e+02;
Matches 16; Conservative 0; Mismatches 2; Indels 0;

Qy      1 TGTAGGAAGCTCTGGGGT 18
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Db      1624 TGGAGGAAGCTTTGGGTG 1641

RESULT 15
US-09-764-855-203
: Sequence 203, Application US/09764855
: Patent No. US20020119919A1
: GENERAL INFORMATION:
: APPLICANT: Rosen et al.
: TITLE OF INVENTION: Nucleic Acids, Proteins, and Antibodies
: FILE REFERENCE: P1110
: CURRENT APPLICATION NUMBER: US/09/764,855
: CURRENT FILING DATE: 2001-01-17
: Prior application data removed - consult PALM or file wrapper
: NUMBER OF SEQ ID NOS: 334
: SOFTWARE: Patentin Ver. 2.0
: SEQ ID NO 203
: LENGTH: 3606
: TYPE: DNA
: ORGANISM: Homo sapiens
US-09-764-855-203

Query Match      77.9%; Score 14.8; DB 10; Length 3606;
Best Local Similarity 88.9%; Pred. No. 1.7e+02;
Matches 16; Conservative 0; Mismatches 2; Indels 0;

Qy      1 TGTAGGAAGCTCTGGGGT 18
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Db      1976 TCGAGGAAGCTTTGGGTG 1993

Search completed: January 21, 2003, 16:29:18
Job time : 37.3056 secs

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RESULT 12
US-09-878-574-4575
; Sequence 4575, Application US/09878574
; Patent No. US20020110548A1
; GENERAL INFORMATION:
; APPLICANT: Byrum, Joseph R.
; APPLICANT: La Rosa, Thomas J.
; APPLICANT: Thompson, Michael D.
; TITLE OF INVENTION: Nucleic Acid Molecules and Other Molecules Associated with
; TITLE OF INVENTION: Plants
; FILE REFERENCE: 38-21(15401)B
; CURRENT APPLICATION NUMBER: US/09/878,574
; CURRENT FILING DATE: 2001-12-21
; PRIOR APPLICATION NUMBER: 09/333,535
; PRIOR FILING DATE: 1999-06-14
; NUMBER OF SEQ ID NOS: 15775
; SEQ ID NO 4575
; LENGTH: 600
; TYPE: DNA
; ORGANISM: Glycine max
; FEATURE:
; NAME/KEY: unsure
; LOCATION: (1)..(600)
; OTHER INFORMATION: unsure at all n locations
; OTHER INFORMATION: Clone ID: LIB3028-033-Q1-BJ-III
US-09-878-574-4575

Query Match 77.9%; Score 14.8; DB 10; Length 600;
Best local similarity 88.9%; Pred. No. 1.5e+02;
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 GTAGGAAGTCTGGGTGC 19
1 | | | | | | | | | | | | | | | |
DB 464 GAGGAAGCCTGGGTGC 481

RESULT 13
US-09-800-729-61
; Sequence 61, Application US/09800729
; Patent No. US20020068319A1
; GENERAL INFORMATION:
; APPLICANT: Ni, et al.
; TITLE OF INVENTION: 32 Human secreted proteins
; FILE REFERENCE: P2044P1
; CURRENT APPLICATION NUMBER: US/09/800,729
; CURRENT FILING DATE: 2001-03-08
; PRIOR APPLICATION NUMBER: PCT/US00/26013
; PRIOR FILING DATE: 2000-09-22
; PRIOR APPLICATION NUMBER: 60/155,709
; PRIOR FILING DATE: 1999-09-24
; NUMBER OF SEQ ID NOS: 217
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 61
; LENGTH: 1499
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-800-729-61

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RESULT 14  
US-09-764-855-202  
; Sequence 202, Application US/09764855  
; Patent NO US20020119919A1

GenCore version 5.1.3  
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UM nucleic - nucleic search, using sw model

Run on: January 21, 2003, 15:09:37 ; Search time 32.7222 Seconds  
(without alignments)  
178.070 Million cell updates/sec

Title: US-09-853-688-36

Perfect score: 19

Sequence: 1 lgtaggaagctctgggtgc 19

Scoring table: IDENTITY\_NUC

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Searched: 441362 seqs, 153338381 residues

Total number of hits satisfying chosen parameters: 882724

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : Issued\_Patents\_NA:\*  
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2: /cgn2.6/ptodata/2/ina/5B.COMB.seq:\*  
3: /cgn2.6/ptodata/2/ina/6A.COMB.seq:\*  
4: /cgn2.6/ptodata/2/ina/6B.COMB.seq:\*  
5: /cgn2.6/ptodata/2/ina/PCTUS.COMB.seq:\*  
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pred. No. is the number of results predicted by chance to have a  
score greater than or equal to the score of the result being printed,  
and is derived by analysis of the total score distribution.

SUMMARIES

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C 3	15.8	83.2	2246	4	US-09-032-742-22
C 4	15.8	83.2	2246	4	US-09-032-742-24
C 5	15.8	83.2	2246	4	US-09-032-742-25
C 6	14.8	77.9	3388	4	US-09-141-206-1
C 7	14.4	75.8	2511	2	US-08-422-699A-8
C 8	14.4	75.8	2511	2	US-08-422-706B-8
C 9	14.4	75.8	2726	2	US-08-422-699A-12
C 10	14.4	75.8	2726	2	US-08-422-706B-12
C 11	14.4	75.8	3182	1	US-08-484-044-11
C 12	14.4	75.8	3323	2	US-08-422-699A-10
C 13	14.4	75.8	3323	2	US-08-422-706B-10
C 14	14.4	75.8	11613	1	US-08-484-044-10
C 15	14.4	75.8	20303	1	US-08-370-975B-6
C 16	14.4	75.8	26764	1	US-08-370-975B-1
C 17	14.4	75.8	72928	3	US-09-009-913-1
C 18	14.2	74.7	94	1	US-08-909-725-1
C 19	14.2	74.7	586	4	US-09-146-969-3
C 20	14.2	74.7	704	4	US-08-998-416-977
C 21	14.2	74.7	747	2	US-08-401-530A-1
C 22	14.2	74.7	747	2	US-08-709-662-1
C 23	14.2	74.7	1878	4	US-09-732-025-1
C 24	14.2	74.7	2291	1	US-07-872-644-5
C 25	14.2	74.7	2291	1	US-08-297-494-5
C 26	14.2	74.7	2291	1	US-08-297-510-5
C 27	14.2	74.7	2291	1	US-08-479-532-5

28	14.2	74.7	2291	1	US-08-455-526-5	Sequence 5, Appli
29	14.2	74.7	2291	1	US-08-455-525-5	Sequence 5, Appli
30	14.2	74.7	2291	3	US-09-139-491-5	Sequence 5, Appli
31	14.2	74.7	2291	5	PCT-US92-03222-5	Sequence 5, Appli
C 32	14.2	74.7	2808	3	US-08-870-126-7	Sequence 7, Appli
C 33	14.2	74.7	2808	4	US-09-445-247-7	Sequence 7, Appli
C 34	14	73.7	3808	2	US-08-916-917-3	Sequence 3, Appli
C 35	14	73.7	3808	2	US-08-972-629-3	Sequence 3, Appli
C 36	14	73.7	3808	2	US-08-972-629-3	Sequence 3, Appli
C 37	14	73.7	3808	2	US-08-972-630-3	Sequence 3, Appli
C 38	14	73.7	3808	2	US-08-672-211-3	Sequence 3, Appli
C 39	14	73.7	3808	3	US-09-225-170-3	Sequence 3, Appli
40	13.8	72.6	588	4	US-09-385-982-128	Sequence 128, App
41	13.8	72.6	1646	4	US-09-221-017B-654	Sequence 654, App
C 42	13.8	72.6	2968	4	US-09-813-819-1	Sequence 1, Appli
C 43	13.8	72.6	2968	4	US-09-920-048-1	Sequence 1, Appli
C 44	13.8	72.6	3022	4	US-09-193-562D-33	Sequence 33, Appli
45	13.8	72.6	11832	2	US-08-416-603-1	Sequence 1, Appli

ALIGNMENTS

RESULT 1  
US-09-032-742-3/c  
; Sequence 3, Application US/09032742  
; Patent No. 6255089  
; GENERAL INFORMATION:  
; APPLICANT: Teitler, Milt  
; APPLICANT: Herrick-Davis, Katharine  
; TITLE OF INVENTION: Constitutively Activated Serotonin  
; TITLE OF INVENTION: Receptors  
; NUMBER OF SEQUENCES: 25  
; CORRESPONDENCE ADDRESS:  
; ADDRESSEE: Laurence Weinberger  
; STREET: 882 S. Matlack Street, Suite 103  
; CITY: West Chester  
; STATE: PA  
; COUNTRY: USA  
; ZIP: 19380-0053  
; COMPUTER READABLE FORM:  
; MEDIUM TYPE: Floppy disk  
; COMPUTER: IBM PC compatible  
; OPERATING SYSTEM: PC-DOS/MS-DOS  
; SOFTWARE: Patentin Release #1.0, version #1.30  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/09/032,742  
; FILING DATE: 27-FEB-1998  
; CLASSIFICATION: 536  
; ATTORNEY/AGENT INFORMATION:  
; NAME: Weinberger, Laurence  
; REGISTRATION NUMBER: 27,965  
; REFERENCE/DOCKET NUMBER: 3086-4  
; TELECOMMUNICATION INFORMATION:  
; TELEPHONE: (610) 431-1703  
; TELEFAX: (610) 431-4181  
; INFORMATION FOR SEQ ID NO: 3:  
; SEQUENCE CHARACTERISTICS:  
; LENGTH: 2246 base pairs  
; TYPE: nucleic acid  
; STRANDEDNESS: single  
; TOPOLOGY: linear  
; MOLECULE TYPE: DNA (genomic)  
US-09-032-742-3

Query Match 83.2%; Score 15.8; DB 4; Length 2246;  
Best Local Similarity 89.5%; Pred. No. 25;  
Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 TGTAGGAAGCTCTGGGTGC 19  
|||||

Db 2096 TGTAGGAAGTCTCGCGTGC 2078

## RESULT 2

US-09-032-742-21/c

Sequence 21, Application US/09032742

Patent No. 6255089

GENERAL INFORMATION:

APPLICANT: Teitler, Milt

APPLICANT: Herrick-Davis, Katharine

APPLICANT: Egan, Christina C.

TITLE OF INVENTION: Constitutively Activated Serotonin

NUMBER OF INVENTIONS: 25

CORRESPONDENCE ADDRESS:

ADDRESSEE: Laurence Weinberger

STREET: 882 S. Matlack Street, Suite 103

STREET: P.O. Box 1663

CITY: West Chester

STATE: PA

COUNTRY: USA

ZIP: 19380-0053

COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk

COMPUTER: IBM PC compatible

OPERATING SYSTEM: PC-DOS/MS-DOS

SOFTWARE: Patent In Release #1.0, Version #1.30

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/09/032,742

FILING DATE: 27-FEB-1998

CLASSIFICATION: 536

ATTORNEY/AGENT INFORMATION:

NAME: Weinberger, Laurence

REGISTRATION NUMBER: 27,965

REFERENCE/DOCKET NUMBER: 3086-4

TELECOMMUNICATION INFORMATION:

TELEPHONE: (610) 431-1703

TELEFAX: (610) 431-4181

INFORMATION FOR SEQ ID NO: 21:

SEQUENCE CHARACTERISTICS:

LENGTH: 2246 base pairs

TYPE: nucleic acid

STRANDEDNESS: single

TOPOLOGY: linear

MOLECULE TYPE: DNA (genomic)

US-09-032-742-21

Query Match 83.2%; Score 15.8; DB 4; Length 2246;

Best Local Similarity 89.5%; Pred. No. 25;

Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 TGTAGGAAGTCTCGCGTGC 19

Db 2096 TGTAGGAAGTCTCGCGTGC 2078

## RESULT 3

US-09-032-742-22/c

Sequence 22, Application US/09032742

Patent No. 6255089

GENERAL INFORMATION:

APPLICANT: Teitler, Milt

APPLICANT: Herrick-Davis, Katharine

APPLICANT: Egan, Christina C.

TITLE OF INVENTION: Constitutively Activated Serotonin

NUMBER OF INVENTIONS: 25

CORRESPONDENCE ADDRESS:

ADDRESSEE: Laurence Weinberger

STREET: 882 S. Matlack Street, Suite 103

STREET: P.O. Box 1663

CITY: West Chester

STATE: PA

1

COUNTRY: USA  
ZIP: 19380-0053

COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk

COMPUTER: IBM PC compatible

OPERATING SYSTEM: PC-DOS/MS-DOS

SOFTWARE: Patent In Release #1.0, Version #1.30

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/09/032,742

FILING DATE: 27-FEB-1998

CLASSIFICATION: 536

ATTORNEY/AGENT INFORMATION:

NAME: Weinberger, Laurence

REGISTRATION NUMBER: 27,965

REFERENCE/DOCKET NUMBER: 3086-4

TELECOMMUNICATION INFORMATION:

TELEPHONE: (610) 431-1703

TELEFAX: (610) 431-4181

INFORMATION FOR SEQ ID NO: 22:

SEQUENCE CHARACTERISTICS:

LENGTH: 2246 base pairs

TYPE: nucleic acid

STRANDEDNESS: single

TOPOLOGY: linear

MOLECULE TYPE: DNA (genomic)

US-09-032-742-22

Query Match 83.2%; Score 15.8; DB 4; Length 2246;

Best Local Similarity 89.5%; Pred. No. 25;

Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 TGTAGGAAGTCTCGCGTGC 19

Db 2096 TGTAGGAAGTCTCGCGTGC 2078

## RESULT 4

US-09-032-742-24/c

Sequence 24, Application US/09032742

Patent No. 6255089

GENERAL INFORMATION:

APPLICANT: Teitler, Milt

APPLICANT: Herrick-Davis, Katharine

APPLICANT: Egan, Christina C.

TITLE OF INVENTION: Constitutively Activated Serotonin

NUMBER OF INVENTIONS: 25

CORRESPONDENCE ADDRESS:

ADDRESSEE: Laurence Weinberger

STREET: 882 S. Matlack Street, Suite 103

STREET: P.O. Box 1663

CITY: West Chester

STATE: PA

COUNTRY: USA

ZIP: 19380-0053

COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk

COMPUTER: IBM PC compatible

OPERATING SYSTEM: PC-DOS/MS-DOS

SOFTWARE: Patent In Release #1.0, Version #1.30

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/09/032,742

FILING DATE: 27-FEB-1998

CLASSIFICATION: 536

ATTORNEY/AGENT INFORMATION:

NAME: Weinberger, Laurence

REGISTRATION NUMBER: 27,965

REFERENCE/DOCKET NUMBER: 3086-4

TELECOMMUNICATION INFORMATION:

TELEPHONE: (610) 431-1703

TELEFAX: (610) 431-4181

INFORMATION FOR SEQ ID NO: 24:

SEQUENCE CHARACTERISTICS:

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; LENGTH: 2246 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
US-09-032-742-24

Query Match      81.2%; Score 15.8; DB 4; Length 2246;
Best Local Similarity 89.5%; Pred. No. 25;
Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 TGTAGGAAGTCTGGGGTGC 19
   |||||
Db 2096 TGTAGGAAGTCTGGCGTGC 2078

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RESULT 5
; Sequence 25, Application US/09032742
; Patent No. 6255089
; GENERAL INFORMATION:
; APPLICANT: Teitler, Milt
; APPLICANT: Herrick-Davis, Katharine
; APPLICANT: Egan, Christina C.
; TITLE OF INVENTION: Constitutively Activated Serotonin
; TITLE OF INVENTION: Receptors
; NUMBER OF SEQUENCES: 25
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Laurence Weinberger
; STREET: 882 S. Matlack Street, Suite 103
; CITY: P.O. Box 1663
; STATE: PA
; COUNTRY: USA
; ZIP: 19380-0053
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/032,742
; FILING DATE: 27-FEB-1998
; CLASSIFICATION: 536
; ATTORNEY/AGENT INFORMATION:
; NAME: Weinberger, Laurence
; REGISTRATION NUMBER: 27,965
; REFERENCE/DOCKET NUMBER: 3086-4
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (610) 431-1703
; TELEFAX: (610) 431-4181
; INFORMATION FOR SEQ ID NO: 25:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 2246 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
US-09-032-742-25

Query Match      83.2%; Score 15.8; DB 4; Length 2246;
Best Local Similarity 89.5%; Pred. No. 25;
Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 TGTAGGAAGTCTGGGGTGC 19
   |||||
Db 2096 TGTAGGAAGTCTGGCGTGC 2078

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RESULT 6
US-09-141-206-1/c
; Sequence 1, Application US/09141206
; Patent No. 6187559

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; GENERAL INFORMATION:
; APPLICANT: Steed, Paul M.
; APPLICANT: Lasala, Daniel J.
; TITLE OF INVENTION: Amino Acid Sequence of Human PLD2A
; Patent No. 6187559
; FILE REFERENCE: 4-30148/PL/CCG1954/R
; CURRENT APPLICATION NUMBER: US/09/141,206
; CURRENT FILING DATE: 1998-08-27
; EARLIER APPLICATION NUMBER: 60/057,802
; EARLIER FILING DATE: 1997-08-28
; NUMBER OF SEQ ID NOS: 8
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 1
; LENGTH: 3388
; TYPE: DNA
; ORGANISM: Human
; FEATURE:
; NAME/KEY: CDS
; LOCATION: (162)..(2963)
; OTHER INFORMATION: Human PLD2
US-09-141-206-1

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Query Match      77.9%; Score 14.8; DB 4; Length 3388;
Best Local Similarity 88.9%; Pred. No. 86;
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 TGTAGGAAGTCTGGGGTG 18
   |||
Db 1663 TGAAGAAGTCTGGGGTG 1646

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RESULT 7
US-08-422-699A-8/c
; Sequence 8, Application US/08422699A
; Patent No. 5955265
; GENERAL INFORMATION:
; APPLICANT: Brook, J. David
; APPLICANT: Housman, David E.
; APPLICANT: Shaw, Duncan J.
; APPLICANT: Harley, Helen G.
; APPLICANT: Johnson, Keith J.
; TITLE OF INVENTION: DNA SEQUENCE ENCODING THE MYOTONIC
; TITLE OF INVENTION: DYSTROPHY GENE AND USES THEREOF
; NUMBER OF SEQUENCES: 14
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Hamilton, Brook, Smith & Reynolds, P.C.
; STREET: Two Militia Drive
; CITY: Lexington
; STATE: Massachusetts
; COUNTRY: US
; ZIP: 02713
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/422,699A
; FILING DATE:
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/422,706
; FILING DATE:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/023,612
; FILING DATE: 26-FEB-1993
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/839,255
; FILING DATE: 20-FEB-1992
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: PCT/US93/01545
; FILING DATE: 19-FEB-1993
; PRIOR APPLICATION DATA:

```

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; APPLICATION NUMBER: PCT/GB93/00253
; FILING DATE: 05-FEB-1993
; PRIOR APPLICATION NUMBER: GB9202485.0
; FILING DATE: 06-FEB-1992
; ATTORNEY/AGENT INFORMATION:
; NAME: Granahan, Patricia
; REGISTRATION NUMBER: 32,227
; REFERENCE/DOCKET NUMBER: MIT-5830A2
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 617-861-6240
; TELEFAX: 617-861-9540
; INFORMATION FOR SEQ ID NO: 8:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 2511 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; FEATURE:
; NAME/KEY: CDS
; LOCATION: 1..1746
;
US-08-422-699A-8

Query Match 75.8%; Score 14.4; DB 2; Length 2511;
Best Local Similarity 93.8%; Pred. No. 1.3e+02;
Matches 15; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 4 AGGAAGTCTGGGGTGC 19
   ||| |||||
DB 590 AGGTAGTCTGGGGTGC 575

RESULT 8
US-08-422-706B-8/c
; Sequence 8, Application US/08422706B
; Patent No. 5977333
; GENERAL INFORMATION:
; APPLICANT: Brook, J. David
; APPLICANT: Housman, David E.
; APPLICANT: Shaw, Duncan J.
; APPLICANT: Harley, Helen G.
; APPLICANT: Johnson, Keith J.
; TITLE OF INVENTION: DNA SEQUENCE ENCODING THE MYOTONIC
; TITLE OF INVENTION: DYSTROPHY GENE AND USES THEREOF
; NUMBER OF SEQUENCES: 14
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Hamilton, Brook, Smith & Reynolds, P.C.
; STREET: Two Militia Drive
; CITY: Lexington
; STATE: Massachusetts
; COUNTRY: US
; ZIP: 02713
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/422,706B
; FILING DATE: 14-APR-1995
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/284,543
; FILING DATE: 08-AUG-1994
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/023,612
; FILING DATE: 26-FEB-1993
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/839,255
; FILING DATE: 20-FEB-1992
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: PCT/US93/01545
```

```
; FILING DATE: 19-FEB-1993
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: PCT/GB93/00253
; FILING DATE: 05-FEB-1993
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: GB9202485.0
; FILING DATE: 06-FEB-1992
; ATTORNEY/AGENT INFORMATION:
; NAME: Granahan, Patricia
; REGISTRATION NUMBER: 32,227
; REFERENCE/DOCKET NUMBER: MIT-5830A2
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 617-861-6240
; TELEFAX: 617-861-9540
; INFORMATION FOR SEQ ID NO: 8:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 2511 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; FEATURE:
; NAME/KEY: CDS
; LOCATION: 1..1746
;
US-08-422-706B-8

Query Match 75.8%; Score 14.4; DB 2; Length 2511;
Best Local Similarity 93.8%; Pred. No. 1.3e+02;
Matches 15; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 4 AGGAAGTCTGGGGTGC 19
   ||| |||||
DB 590 AGGTAGTCTGGGGTGC 575

RESULT 9
US-08-422-699A-12/c
; Sequence 12, Application US/08422699A
; Patent No. 5955265
; GENERAL INFORMATION:
; APPLICANT: Brook, J. David
; APPLICANT: Housman, David E.
; APPLICANT: Shaw, Duncan J.
; APPLICANT: Harley, Helen G.
; APPLICANT: Johnson, Keith J.
; TITLE OF INVENTION: DNA SEQUENCE ENCODING THE MYOTONIC
; TITLE OF INVENTION: DYSTROPHY GENE AND USES THEREOF
; NUMBER OF SEQUENCES: 14
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Hamilton, Brook, Smith & Reynolds, P.C.
; STREET: Two Militia Drive
; CITY: Lexington
; STATE: Massachusetts
; COUNTRY: US
; ZIP: 02713
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/422,699A
; FILING DATE:
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/422,706
; FILING DATE:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/023,612
; FILING DATE: 26-FEB-1993
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/839,255
; FILING DATE: 20-FEB-1992
```

```

; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: PCT/US93/01545
; FILING DATE: 19-FEB-1993
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: PCT/GB93/00253
; FILING DATE: 05-FEB-1993
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: GB9202485.0
; FILING DATE: 06-FEB-1992
; ATTORNEY/AGENT INFORMATION:
; NAME: Granahan, Patricia
; REGISTRATION NUMBER: 32,227
; REFERENCE/DOCKET NUMBER: MIT-5830A2
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 617-861-6240
; TELEFAX: 617-861-9540
; INFORMATION FOR SEQ ID NO: 12:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 2726 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; US-08-422-699A-12

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```

Query Match 75.8%; Score 14.4; DB 2; Length 2726;
Best Local Similarity 93.8%; Pred. No. 1.3e+02;
Matches 15; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

```

```

QY 4 AGGAAGTCTGGGTGC 19
    ||| |||||
DB 840 AGGTAGTCTGGGTGC 825

```

```

RESULT 10
US-08-422-706B-12/C
; Sequence 12, Application US/08422706B
; Patent No. 5977333
; GENERAL INFORMATION:
; APPLICANT: Brook, J. David
; APPLICANT: Housman, David E.
; APPLICANT: Shaw, Duncan J.
; APPLICANT: Harley, Helen G.
; APPLICANT: Johnson, Keith J.
; TITLE OF INVENTION: DNA SEQUENCE ENCODING THE MYOTONIC
; TITLE OF INVENTION: DYSTROPHY GENE AND USES THEREOF
; NUMBER OF SEQUENCES: 14
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Hamilton, Brook, Smith & Reynolds, P.C.
; STREET: Two Militia Drive
; CITY: Lexington
; STATE: Massachusetts
; COUNTRY: US
; ZIP: 02713
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/422,706B
; FILING DATE: 14-APR-1995
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/284,543
; FILING DATE: 08-AUG-1994
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/023,612
; FILING DATE: 26-FEB-1993
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/839,255
; FILING DATE: 20-FEB-1992
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: PCT/US93/01545

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; FILING DATE: 19-FEB-1993
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: PCT/GB93/00253
; FILING DATE: 05-FEB-1993
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: GB9202485.0
; FILING DATE: 06-FEB-1992
; ATTORNEY/AGENT INFORMATION:
; NAME: Granahan, Patricia
; REGISTRATION NUMBER: 32,227
; REFERENCE/DOCKET NUMBER: MIT-5830A2
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 617-861-6240
; TELEFAX: 617-861-9540
; INFORMATION FOR SEQ ID NO: 12:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 2726 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; US-08-422-706B-12

```

```

Query Match 75.8%; Score 14.4; DB 2; Length 2726;
Best Local Similarity 93.8%; Pred. No. 1.3e+02;
Matches 15; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

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QY 4 AGGAAGTCTGGGTGC 19
    ||| |||||
DB 840 AGGTAGTCTGGGTGC 825

```

```

RESULT 11
US-08-484-044-11/c
; Sequence 11, Application US/08484044
; Patent No. 5552282
; GENERAL INFORMATION:
; APPLICANT: Caskey, C. T.
; APPLICANT: Fu, Ying-Hui
; APPLICANT: Friedman, David L.
; APPLICANT: Pizzuti, Antonio
; APPLICANT: Fenwick, Raymond G.
; TITLE OF INVENTION: Diagnosis of Myotonic Muscular Dystrophy
; NUMBER OF SEQUENCES: 13
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Fulbright & Jaworski, L.L.P.
; STREET: 1301 McKinney, Suite 5100
; CITY: Houston
; STATE: Texas
; COUNTRY: U.S.A.
; ZIP: 77010-3095
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/484,044
; FILING DATE:
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/019,940
; FILING DATE: 19-FEB-1993
; ATTORNEY/AGENT INFORMATION:
; NAME: Paul, Thomas D.
; REGISTRATION NUMBER: 32,714
; REFERENCE/DOCKET NUMBER: D-5443
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 713/651-5325
; TELEFAX: 713/651-5246
; TELEX: 762829
; INFORMATION FOR SEQ ID NO: 11:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 3182 base pairs

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TELECOMMUNICATION INFORMATION:  
TELEPHONE: 617-861-6240  
TELEFAX: 617-861-9540  
INFORMATION FOR SEQ ID NO: 10:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 3323 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: double  
TOPOLOGY: linear  
MOLECULE TYPE: DNA (genomic)  
FEATURE:  
NAME/KEY: misc.difference  
LOCATION: replace(518..3323, "")  
OTHER INFORMATION: /standard\_name= "cDNA 41"  
FEATURE:  
NAME/KEY: misc.difference  
LOCATION: replace(769..3323, "")  
OTHER INFORMATION: /standard\_name= "cDNA 28"  
US-08-422-706B-10

Query Match 75.8%; Score 14.4; DB 2; Length 3323;  
Best Local Similarity 93.8%; Pred. No. 1.4e+02;  
Matches 15; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 4 AGGAAGTCTGGGGTGC 19  
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DB 1375 AGGTAGTCTGGGGTGC 1360

## RESULT 14

US-08-484-044-10/c  
Sequence 10, Application US/08484044  
Patent No. 5552282  
GENERAL INFORMATION:  
APPLICANT: Caskey, C. T.  
APPLICANT: Fu, Ying-Hui  
APPLICANT: Friedman, David L.  
APPLICANT: Pizzulli, Antonio  
APPLICANT: Fenwick, Raymond G.  
TITLE OF INVENTION: Diagnosis of Myotonic Muscular Dystrophy  
NUMBER OF SEQUENCES: 13  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Fulbright & Jaworski, L.L.P.  
STREET: 1301 McKinney, Suite 5100  
CITY: Houston  
STATE: Texas  
COUNTRY: U.S.A.  
ZIP: 77010-3095  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: PatentIn Release #1.0, Version #1.25  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/484,044  
FILING DATE:  
CLASSIFICATION: 435  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/019,940  
FILING DATE: 19-FEB-1993  
ATTORNEY/AGENT INFORMATION:  
NAME: Paul, Thomas D.  
REGISTRATION NUMBER: 32,714  
REFERENCE/DOCKET NUMBER: D-5443  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: 713/651-5325  
TELEFAX: 713/651-5246  
TELEX: 762829  
INFORMATION FOR SEQ ID NO: 10:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 11613 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: double

TOPOLOGY: linear  
MOLECULE TYPE: DNA (genomic)  
US-08-484-044-10

Query Match 75.8%; Score 14.4; DB 1; Length 11613;  
Best Local Similarity 93.8%; Pred. No. 1.6e+02;  
Matches 15; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 4 AGGAAGTCTGGGGTGC 19

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DB 4077 AGGTAGTCTGGGGTGC 4062

## RESULT 15

US-08-370-975B-6/c  
Sequence 6, Application US/08370975B  
Patent No. 5622851  
GENERAL INFORMATION:  
APPLICANT: Maley, Frank  
APPLICANT: Maley, Gladys F.  
APPLICANT: Weiner, Karen X.B.  
TITLE OF INVENTION: Human Deoxycytidylate Deaminase Gene  
NUMBER OF SEQUENCES: 14  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Nixon, Hargrave, Devans & Doyle  
STREET: Clinton Square, P.O. Box 1051  
CITY: Rochester  
STATE: New York  
COUNTRY: USA  
ZIP: 14603  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: PatentIn Release #1.0, Version #1.25  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/370,975B  
FILING DATE: 10-JAN-1995  
CLASSIFICATION: 435  
ATTORNEY/AGENT INFORMATION:  
NAME: Timlan, Susan J.  
REGISTRATION NUMBER: 34,103  
REFERENCE/DOCKET NUMBER: 20894/80  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: (716)263-1636  
TELEFAX: (716)263-1600  
INFORMATION FOR SEQ ID NO: 6:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 20303 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: single  
TOPOLOGY: linear  
MOLECULE TYPE: DNA (genomic)  
POSITION IN GENOME:  
CHROMOSOME/SEGMENT: 4q35  
US-08-370-975B-6

Query Match 75.8%; Score 14.4; DB 1; Length 20303;  
Best Local Similarity 93.8%; Pred. No. 1.7e+02;  
Matches 15; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 3 TAGGAAGTCTGGGGTG 18

||| |||||

DB 2205 TGGGAAGTCTGGGGTG 2190

Search completed: January 21, 2003, 16:30:42  
Job time : 36.7222 secs